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Physical Properties of Bedding Materials Determine the Marble Burying Behavior of Mice (C57BL/6J)

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Abstract: The marble burying test is a simple behavioral psychopharmacology task in which rodents, especially rats and mice, are exposed to glass marbles laying on thick bedding materials. This test has been widely used to assess the effects of a variety of psychoactive drugs. Although the marble burying test clearly reflects drug efficacy, many aspects of the behavior itself remain unclear. In the present study, we assessed the effects of different bedding materials and the number of glass marbles presented on the marble burying behavior of mice (C57BL/6J). Six (bedding materials) × three (number of glass marbles) factorial experiments revealed that the number of marbles buried correlates closely with the quality of the bedding material. Our results provide a basic understanding of certain factors that govern marble burying behavior, which may be applicable to behavioral sciences including psychopharmacology.

Keywords: Marble burying, defensive burying, digging, bedding materials, mice.

INTRODUCTION

Defensive burying is a widely observed behavior in rodents [1-8]. Rats and mice eagerly bury noxious things, such as bad tasting and/or smelling food [1-3], shock prod [4], and unusual objects like glass marbles [2], under their bedding materials. This behavior is thought to be an expression of defensiveness, some forms of anxiety, or compulsiveness [5]. When rats and/or mice are placed in a cage in which glass marbles are laying on sawdust, they immediately and eagerly begin burying the marbles [2]. The marble burying behavior is quite easy to observe and is sensitive to anxiolytic and/or anxiogenic compounds. Therefore, this behavior is often used to assess the properties of psychoactive drugs [6-11].

Certain unique properties of marble burying behavior have been revealed. Animals bury glass marbles even though they are not harmful. Marble burying behavior does not show any habituation after repeated trials [6,12-14], indicating that this behavior is not likely to be affected by learning processes or cognitive function. Furthermore, marble burying behavior does not differ by gender [12]. Although marble burying behavior has been well characterized, the factors governing the burying behavior itself remain unclear. It was recently reported that the marble burying test simply reflects digging and other related behaviors [14-17], calling into question the validity of this behavior for assessing anxiety, defensiveness, compulsiveness, and so on. However, marble burying behavior correlates well with both serotonin (5-HT) transporter (SERT) binding of selective 5-HT reuptake inhibitors (SSRIs) and extracellular 5-HT levels in brains treated with SSRIs [18,19], and thus the behavior may reflect a relation between 5-HT neuronal

activity and anxiety, defensiveness, compulsiveness. Therefore, the marble burying test is often used to assess the efficacy of drugs such as SSRIs.

The number of marbles an animal buries is determined by certain factors such as the type of bedding material and the number of marbles used. However, these aspects have not been well studied. For example, many studies have used "saw dust" as the bedding material to study burying behavior because it is lightweight, allowing animals to easily bury the marbles. This feature of "saw dust" might prompt digging, however, leading to a misunderstanding of the emotional state of the animals and/or drug effects. Thus, knowledge of the effects of different bedding materials is required to properly assess burying behavior.

The aim of this study was to assess the effects of bedding material type and number of glass marbles on marble burying test results. Two-factorial within-subject analysis revealed significant effects of bedding materials on the marble burying behavior of C57BL/6J mice.

MATERIALS AND METHODOLOGY

Animals

Seventeen male C57BL/6J mice were used as subjects. Mice were bred in the animal facility of RIKEN BSI (originally purchased from JCL, Inc. Tokyo, Japan, F2 generation) and were about 10 weeks of age at the beginning of the experiment. The breeding room was air-conditioned (22 ± 1°C, 50-60% humidity) with a 12h:12h light-dark cycle (lights on at 0800). Mice were individually housed beginning one week prior to the experiment, and food and water were freely available except during experimentation. All experiments were conducted in the breeding room during the light cycle between 1300 and 1700.

All animal experiments in this study were performed in strict accordance with the guidelines of The Institute of Physical and Chemical Research (RIKEN), Brain Science

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Institute (BSI), and were approved by the Animal Investigation Committee of the Institute.

Apparatus

All experiments were conducted in clear plastic caging with filter tops (Allentown, PA, USA: 29 × 19.5 × 13.5 cm (H)). Glass marbles were clear, green in color and 1.5 cm in diameter. A spontaneous activity monitor (AB System: Neuroscience, Tokyo, Japan) was used to measure activity in the cage during both habituation and test periods.

Bedding Materials

Six types of commercially available bedding materials were used: Soft-chip and Paper-clean (SLC, Shizuoka, Japan), TEK-Fresh (Harlan, IN, USA), Cellu-dri and ALPHA-dri (Shepherd Specialty Papers, TN, USA), and GreenTru (GREENPRODUCTS, IA, USA) (Fig. 1). All bedding materials were autoclaved before use.

Spontaneous Activity Monitoring

The ceiling of each compartment of the breeding rack was equipped with a pyroelectric sensor to monitor the movements of mice throughout the experiment. Data were collected and analyzed with a PC equipped with Windows and other commercially available software (AB System). Activity counts represent the number of active time bin (approximately 0.20 - 0.25 sec each) in which spontaneous activity including locomotor activity, rearing and other activities such as stereotypic movements were detected.

Marble Burying Test

A modified procedure based on Yamada *et al.* [20] was employed. Mice were placed individually in plastic cages with the designated bedding material for 30 min (habituation period) and then placed into waiting cages. Ten, twenty or forty glass marbles were then evenly spaced 3 to 7 cm apart on a 4-cm layer of bedding material in the habituation cages. Mice were then reintroduced into the same cage in which they had been habituated. After 30 min, the marble burying period was terminated by removing the mice, and the number of marbles that were more than two-thirds covered with bedding material was counted. Whether a marble was buried was established by two experimenters and was

confirmed by manual post-experimental assessment of digital photographs on a PC. After each trial, cages and bedding materials were replaced with fresh ones, and glass marbles were washed with water, dried with a paper towel, and left to return to room temperature.

The marble burying test was repeated twenty times. The first, tenth and the last tests were calibration trials, in which mice were exposed to twenty glass marbles on TEK-Fresh, the bedding material used in our breeding facility. In the second to ninth and eleventh to nineteenth trials each mouse was assigned to one of 17 designated conditions—number of glass marbles (10, 20, or 40) and bedding materials (Soft-chip, TEK-Fresh, Paper-clean, Cellu-dri, ALPHA-dri, or GreenTru). The order of condition for each mouse was counterbalanced using *Latin square design*. Each trial was separated by two to three days. On days between trials, mice were individually housed in a cage with a 2-cm layer of TEK-Fresh.

Statistical Analysis

Statistical analyses were performed by the SPSS™ statistical package (ver 16.0). The repeated measure ANOVA and one-way within-subject ANOVA were used. *Post-hoc* analysis was conducted using Dunnett's *t*-test or Tukey's HSD test. The correlation between the density of the bedding materials and the number of buried marbles was analyzed using Pearson's correlation coefficient. Statistical significance was set at $p < 0.05$.

RESULTS

Effects of Repeated Trials on Spontaneous Activity and Marble Burying Behavior

Because many trials were conducted in this study, we assessed the effects of repeated trials both on spontaneous activity and marble burying behavior (Fig. 2). Spontaneous activity increased linearly over the trials, and one-way ANOVA revealed a statistically significant main effect of trial number ($F(2,32) = 10.2$, $p < 0.001$, Fig. 2A). In contrast, the number of buried marbles did not differ significantly with trial number ($F(2,32) = 2.16$, Fig. 2B), consistent with previous studies [6,12,13].

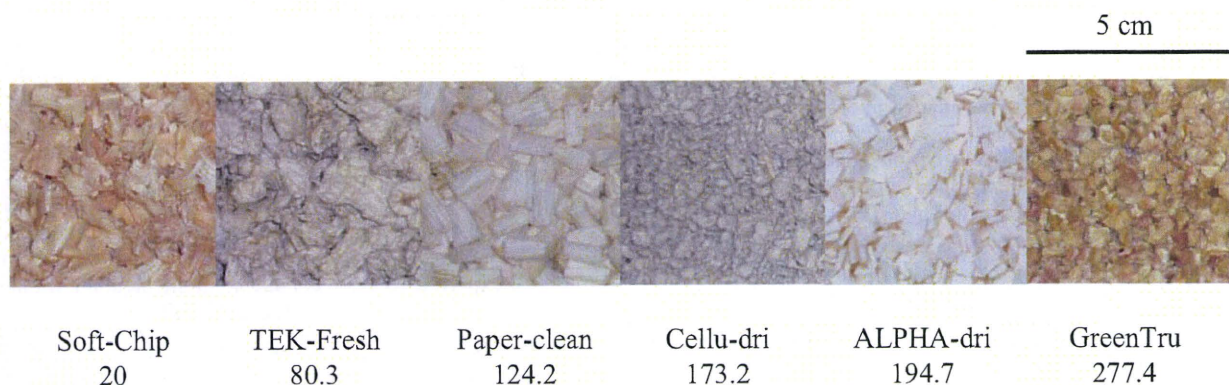
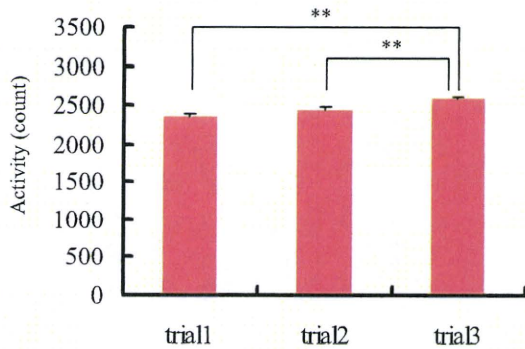


Fig. (1). Bedding materials used in this study. Numbers under the bedding material names indicate the density (g/L).

A Spontaneous activity



B Marble burying

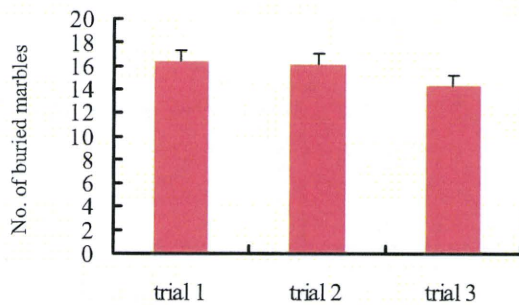


Fig. (2). Spontaneous activity increases with repeated trials, but the number of marbles buried does not. (A) spontaneous activity, (B) number of marbles buried. Number of marbles used was twenty in each trial. Data represent mean ± SEM. **: $p < 0.01$.

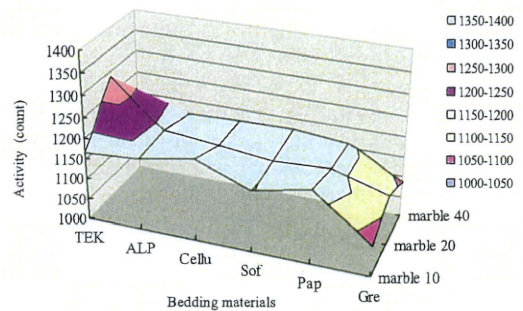
Spontaneous Activity is Affected by Bedding Material

It is possible that the spontaneous activity level of mice could affect marble burying behavior and might be related to properties of the bedding material. Therefore, we assessed the spontaneous activity of mice during both habituation and the marble burying period (Fig. 3). Activity was clearly affected by the type of bedding material. During the habituation period (Fig. 3A), one-way ANOVA revealed a significant main effect of bedding material in 10-, 20-, and 40-marble trials ($F(5,80) = 6.03, p < 0.0001$; $F(5,80) = 7.01, p < 0.0001$; $F(5,80) = 4.37, p < 0.01$, respectively). During the marble burying period (Fig. 3B), one-way ANOVA revealed a significant main effect of bedding material in the 20- and 40-marble trials ($F(5,80) = 2.52, p < 0.05$; $F(5,80) = 3.11, p < 0.05$, respectively). Throughout the experiment, mice exhibited the highest activity in the TEK-Fresh trials and showed the lowest activity in GreenTru trials. These results strongly suggest that spontaneous activity of mice is affected by bedding material properties.

Low-Density Bedding Increase Marble Burying Behavior in Mice

Results of the marble burying test are summarized in Fig. (4). Mice buried many glass marbles when the density of

A Habituation period



B Marble burying period

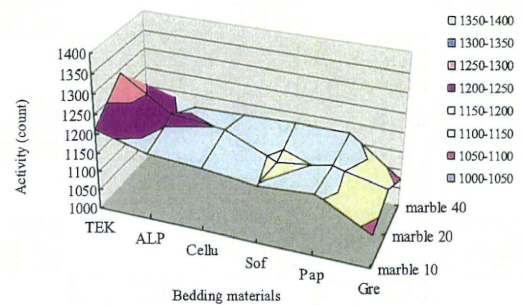


Fig. (3). Mean spontaneous activity during habituation and the marble burying period. (A) habituation period; (B) marble burying period. The terms 'marble 10', 'marble 20', and 'marble 40' indicate marble burying test trials with 10, 20, and 40 marbles, respectively. TEK: TEK-Fresh, ALP: ALPHA-dri, Cellu: Cellu-dri, Sof: Soft-chip, Pap: Paper-clean, Gre: GreenTru. Statistics are presented in the text.

bedding material was low, for example in Soft-chip (20 g/L) and TEK-Fresh (80.3 g/L) bedding. In contrast, they buried only a few marbles when the density of the bedding material was high, for example in ALPHA-dri (194.7 g/L) and GreenTru (277.4 g/L) bedding. In Paper-clean (124.2 g/L) and Cellu-dri (173.2 g/L) bedding, which are moderately dense, the marble burying behavior was moderate. Furthermore, the effect of the number of glass marbles on the burying behavior was salient. In Soft-chip, TEK-Fresh, Cellu-dri and Paper-clean (partially) trials, the number of buried marbles increased linearly with the number of marbles in the cage. The number of buried marbles did not show a linear increment in the ALPHA-dri and GreenTru trials, however. Fig. (5) shows the negative correlation between the number of buried marbles and the density of bedding materials (10 marbles (Fig. 5A): $r = -0.973, p < 0.01$; 20 marbles (Fig. 5B): $r = -0.962, p < 0.01$; 40 marbles (Fig. 5C): $r = -0.941, p < 0.01$, respectively). These results indicate that bedding material density strongly affects the number of marbles buried by mice. In addition, bedding material density may also affect how the mice bury the marbles. Representative photos of burying behaviors in each of the bedding materials are shown in Fig. (6). In Soft-chip and TEK-Fresh, mice completely buried the marbles under

the low-density bedding material. In Cellu-dri, ALPHA-dri and GreenTru, however, the mice only partially buried the marbles. The texture, as well as density, of the bedding material may also affect burying behavior. Mice buried more marbles in bedding materials with a soft texture (Soft-chip, TEK-Fresh, Paper-clean, Cellu-dri) than in bedding with a hard texture (ALPHA-dri, GreenTru). We replicated experiments several times with a somewhat smaller design and obtained almost identical results (data not shown). These results suggest that bedding material properties may affect how mice bury marbles.

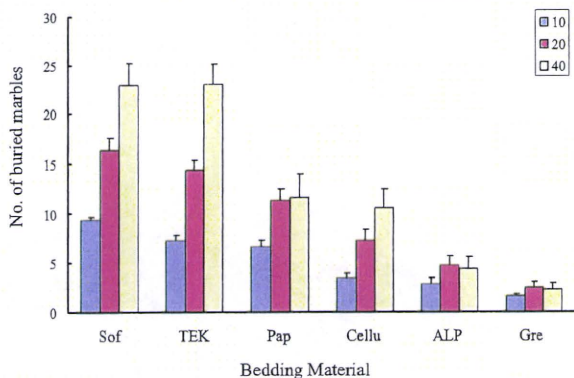


Fig. (4). Marble burying changes with the type of bedding and number of marbles presented. Sof: Soft-chip, TEK: TEK-Fresh, Pap: Paper-clean, Cellu: Cellu-dri, ALP: ALPHA-dri, Gre: GreenTru. 10, 20, and 40 represent trials with 10, 20, and 40 marbles, respectively. Data represent mean ± SEM. Statistics are presented in the text.

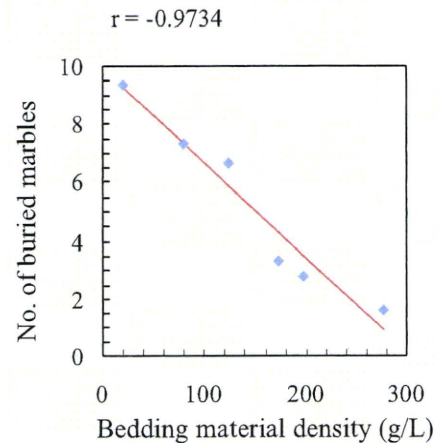
Correlation Between Spontaneous Activity and Number of Marbles Buried

As described above, spontaneous activity may affect marble burying behavior. Therefore, we examined the correlation between the spontaneous activity score and the number of marbles buried (Fig. 7). In both 10- and 20-marble trials, correlation coefficients were not statistically significant, contrary, in the 40-marble trials, the correlation coefficient was statistically significant ($r = 0.20, p < 0.05$). Although there is a significant correlation between spontaneous activity and number of marbles, the small effect size for this measure suggests that spontaneous activity does not play a crucial role for marble burying behavior.

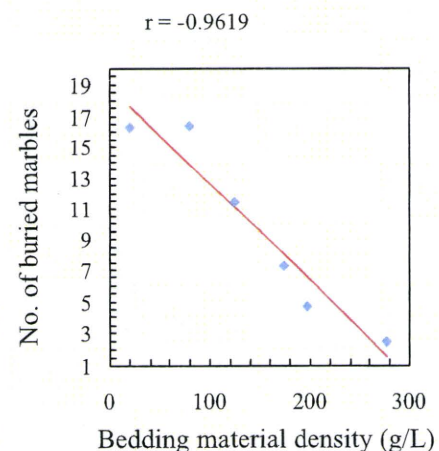
DISCUSSION

In this study, we demonstrated that the physical properties of bedding material and the number of glass marbles in the cage significantly affect the marble burying behavior of mice. The number of marbles buried depended on bedding material density and texture. Mice buried more marbles in low-density, soft bedding than in high-density, hard bedding (Figs. 1, 4). Furthermore, the number of marbles buried correlated with the number of marbles used for the test when the bedding was not high-density and/or hard in texture (Fig. 4). Thus, the properties of bedding, rather than the number of marbles used, seems to be the principal determining factor governing marble burying behavior.

A 10 marbles



B 20 marbles



C 40 marbles

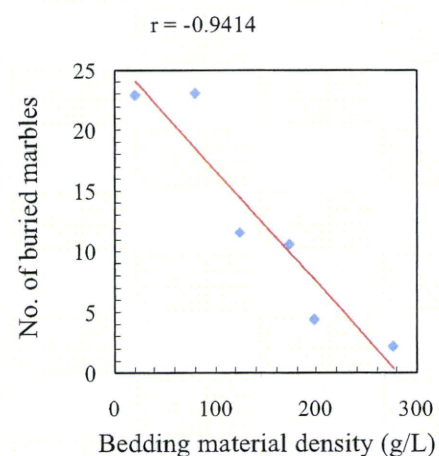


Fig. (5). Correlation between the number of marbles buried and bedding material density. (A) trials with 10 marbles; (B) trials with 20 marbles; (C) trials with 40 marbles. Red lines represent the linear regression.

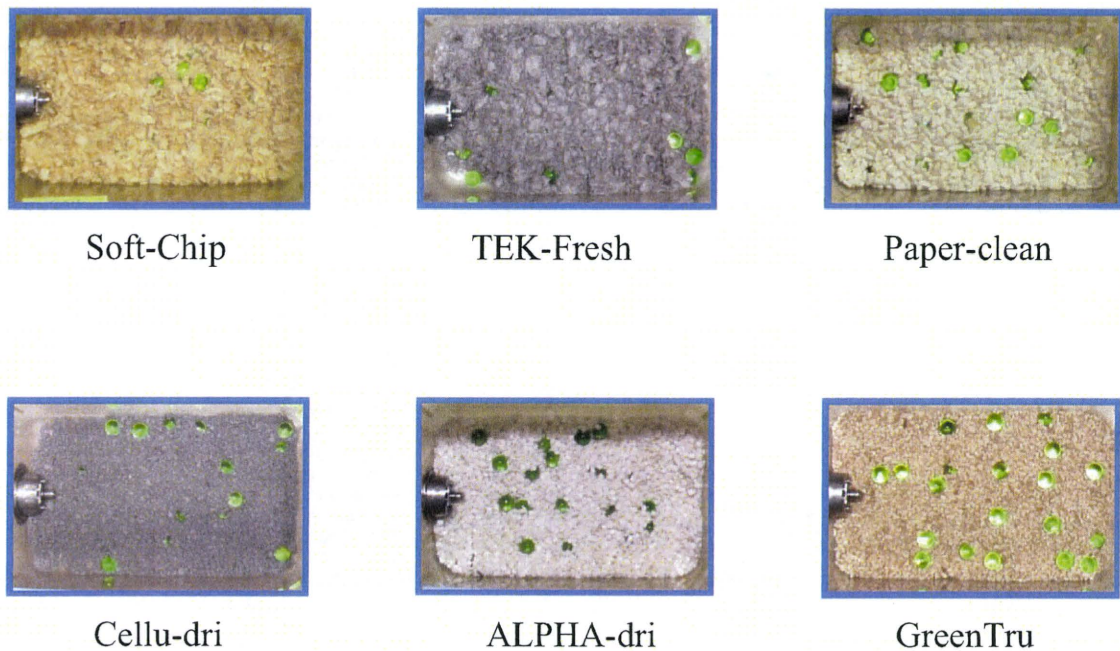


Fig. (6). Representative marble burying behavior in each bedding material.

Bedding material also affected spontaneous activity of mice both quantitatively and qualitatively. Animals were most active in the TEK-Fresh trial and were least active in the GreenTru trial (Fig. 3). The highest activity in the TEK-Fresh trial might be attributed to the 'familiar environment' not to the physical property of the bedding, because TEK-Fresh is used in our breeding facility. However, this possibility might be excluded in this study because mice show high activity when they are exposed to fresh bedding, showing it is not 'familiar'. Activity also differed qualitatively with bedding type. Mice eagerly dug the layer of bedding in TEK-Fresh and Soft-chip trials, but rarely dug in GreenTru trials (data not shown). Certain studies have suggested that marble burying behavior correlates strongly with digging behavior [14-16], and some marbles might be covered with bedding as an indirect consequence of digging. Thus, marble burying behavior, at least in trials using low-density bedding materials, consists of two qualitatively different behaviors, burying and digging. These mixed behaviors may confound the results of studies assessing drugs effects.

Because animals did not dig much in ALPHA-dri and GreenTru trials, the small number of marbles buried in these trials (Fig. 4) might reflect "real burying behavior", raising a fundamental question about the marble burying test. In TEK-Fresh and Soft-chip trials, a majority of marbles might have been buried indirectly by the digging behavior. Previous work has suggested that the marble burying test simply measures digging behavior [14-15]. Is marble burying

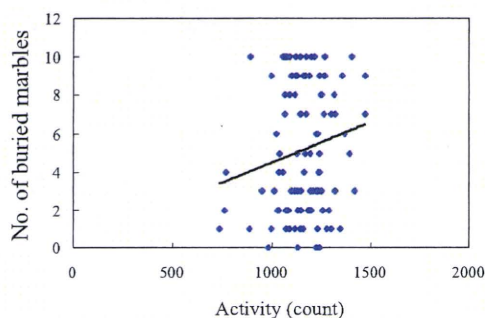
behavior equal to the digging behavior? Or is the contribution of burying to marble burying behavior so small? In this way, marble burying test has some theoretical confusion. Thus, to elucidate this confusion we added some experiments in which the digging behavior in the habituation trial was measured using TEK-Fresh, Soft-chip and GreenTru. The correlation coefficient between the digging duration and the number of buried marbles was statistically significant in Soft-chip experiment ($n=5$, $r=0.897$, $p<0.05$), contrary, not significant in TEK-Fresh and GreenTru experiments ($n=6$, $r=-0.01$, n.s.; $n=7$, $r=0.208$, n.s., respectively). These results suggest that a selection of bedding materials (e.g. TEK-Fresh and/or GreenTru) may be able to dissect out marble burying behavior.

Regardless, the marble burying test is still quite useful for assessing psychotropic drugs [6-11]. For example, in this study, we repeated the marble burying test 20 times for each set of mice, and the number of buried marbles did not change (Fig. 2B). This lack of habituation provides reliability and validity to experiments having a within-subject design. The marble burying test may have other advantages in psychopharmacological studies, but further study is required to elucidate these advantages.

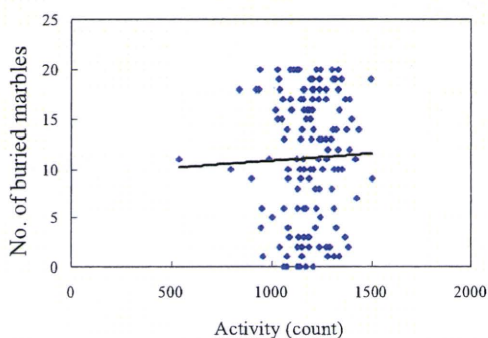
ACKNOWLEDGEMENTS

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A 10 marble trials



B 20 marble trials



C 40 marble trials

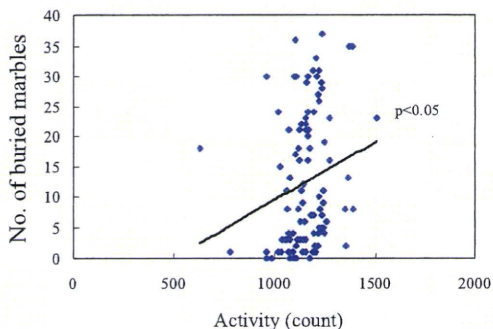


Fig. (7). Spontaneous activity and the number of marbles buried are significantly correlated only for 40-marble trials ($r = 0.20$, $p < 0.05$).

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Social isolation stress induces ATF-7 phosphorylation and impairs silencing of the 5-HT 5B receptor gene

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Many symptoms induced by isolation rearing of rodents may be relevant to neuropsychiatric disorders, including depression. However, identities of transcription factors that regulate gene expression in response to chronic social isolation stress remain elusive. The transcription factor ATF-7 is structurally related to ATF-2, which is activated by various stresses, including inflammatory cytokines. Here, we report that *Atf-7*-deficient mice exhibit abnormal behaviours and increased 5-HT receptor 5B (*Htr5b*) mRNA levels in the dorsal raphe nuclei. ATF-7 silences the transcription of *Htr5b* by directly binding to its 5'-regulatory region, and mediates histone H3-K9 trimethylation via interaction with the ESET histone methyltransferase. Isolation-reared wild-type (WT) mice exhibit abnormal behaviours that resemble those of *Atf-7*-deficient mice. Upon social isolation stress, ATF-7 in the dorsal raphe nucleus is phosphorylated via p38 and is released from the *Htr5b* promoter, leading to the upregulation of *Htr5b*. Thus, ATF-7 may have a critical role in gene expression induced by social isolation stress.

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Introduction

ATF-7 (originally called ATF α) is structurally related to ATF-2 (Hai *et al*, 1989; Maekawa *et al*, 1989; Gaire *et al*, 1990), a member of the ATF-CREB family of transcription factors. ATF-2, ATF-7, and CRE-BPa (Nomura *et al*, 1993) form a subfamily in the ATF-CREB family. Each of these three factors contains a transcription-activation domain consisting of a metal-finger structure and stress-activated protein kinase (SAPK) phosphorylation sites, and a b-ZIP type DNA-binding domain. Various stresses, including inflammatory cytokines, activate SAPKs such as p38 and JNK (Davis, 2000), which then phosphorylate ATF-2 and activate its *trans*-activating capacity (Gupta *et al*, 1995; Livingstone *et al*, 1997; van Dam *et al*, 1997). ATF-7 is also phosphorylated by p38, but not by JNK (De Graeve *et al*, 1999). ATF-2 and ATF-7 can form homodimers or heterodimers with Jun and bind to cAMP response element (CRE) (5'-TGACGTC-3') (Chatton *et al*, 1994).

ATF-7 binds to mouse ATF α -associated modulator (mAM) which is a component of the ESET complex (De Graeve *et al*, 2000; Wang *et al*, 2003). As ESET is a histone methyltransferase (HMTase) that converts lysine 9 of histone H3 (H3-K9) from the dimethyl to the trimethyl form, therefore ATF-7 is thought to support gene silencing by inducing histone H3-K9 trimethylation. Two reports have suggested a role for ATF-2 family transcription factors in epigenetic gene silencing. The yeast homologue of ATF-2, Atf1, contributes to heterochromatin formation independently of the RNAi machinery (Jia *et al*, 2004). Vertebrate ATF-2 also interacts with the histone variant macroH2A, which is enriched in the inactive X chromosome in female mammalian cells and functions to maintain gene silencing (Agelopoulos and Thanos, 2006).

Both ATF-7 and ATF-2 are ubiquitously expressed in various tissues, including the brain (Takeda *et al*, 1991; Goetz *et al*, 1996). *Atf-2* null mice die immediately after birth because of defects in respiration, which appear to be caused by impaired proliferation of cytotrophoblasts in the placenta (Maekawa *et al*, 1999). *Atf-2* heterozygotes are highly prone to mammary tumours in which the expression levels of *Maspin*, a tumour suppressor, and *Gadd45 α* , which is induced by hypoxic stress, are decreased (Maekawa *et al*, 2007). Both these genes encode the regulators of apoptosis, suggesting that defects in the apoptotic machinery are linked to the occurrence of mammary tumours. In contrast, the physiological role of ATF-7 is unknown, although the *Atf-2* and *Atf-7* double mutant exhibits embryonic lethality with abnormalities in the developing liver and heart (Breitwieser *et al*, 2007).

Human neuropsychiatric disorders, such as depression, have multiple-risk factors, including environmental and genetic factors. A loss of social contact is one environmental factor that appears to be linked to both the onset and relapse

of depression (Paykel *et al*, 1980). Long-term social isolation of rodents after weaning provides a model to study the behavioural consequences of loss of social interactions. Many of the symptoms induced by isolation rearing may be relevant to neuropsychiatric disorders (Rodgers and Cole, 1993). Isolated animals are aggressive and exhibit anxiety-like behaviours and increased locomotor activity (Rodgers and Cole, 1993; Blanchard *et al*, 2001). One of the typical abnormal behaviour in isolation-reared mice is a deficit in pre-pulse inhibition (PPI) of the acoustic startle response (Wilkinson *et al*, 1994). In fact, isolation-induced disruption of PPI has been used as a disease model in screening antipsychotic drugs. In animal studies, isolation stress changes the activity of brain neurotransmitters (Blanc *et al*, 1980; Blanchard *et al*, 2001). In the case of acute stress, several transcription factors, including c-Fos and corticosteroid receptors, are activated and modulate multiple target genes (Kaufer *et al*, 1998). However, the transcription factors that are activated and the regulation of gene expression patterns in response to a chronic stress, such as social isolation stress, remain elusive. In addition, as the effect of social isolation stress on behaviour is long-lived, this stress may cause epigenetic changes. However, the mechanism by which epigenetic change is caused by isolation stress remains unknown.

In this study, we have demonstrated that *Atf-7*-deficient (*Atf-7*^{-/-}) mice exhibit abnormal behaviours reminiscent of isolation-reared wild-type (WT) mice. Social isolation stress induced the phosphorylation of ATF-7 and p38 in the dorsal raphe nuclei, as well as a release of ATF-7 from the promoter of the 5-HT receptor 5B (*Htr5b*) gene, leading to an impaired silencing of this gene.

Results

Abnormal behaviours of *Atf-7*^{-/-} mice

We generated *Atf-7*^{-/-} mice (Supplementary Figure S1), and, under pathogen-free conditions, *Atf-7*^{-/-} mice appeared healthy until at least 12 months of age. As *Atf-7* mRNA is expressed at relatively high levels in parts of the brain (Goetz *et al*, 1996), we examined various behaviours originally using WT and *Atf-7*^{-/-} littermate mice with a mixed CBA (25%) × C57BL/6 (75%) genetic background, and later using C57BL/6 congenic mice. In the marble-burying test, which is used to examine anxiety-related behaviours (Spooren *et al*, 2000), *Atf-7*^{-/-} mice exhibited increased marble-burying behaviour compared with WT mice (Figure 1A and Supplementary Figure S2A). In other tests of anxiety-related behaviours, such as the amount of time spent in the centre of an open-field and the elevated plus-maze test (Spooren *et al*, 2000), there was no significant difference between *Atf-7*^{-/-} and WT mice (Figure 1B and C and Supplementary Figure S2B). *Atf-7*^{-/-} mice did exhibit a significant increase in the startle response to a pulse-alone stimulus (Figure 1D and Supplementary Figure S2C). PPI, in which the startle reflex response is attenuated by a pre-pulse, is an important measure of sensorimotor gating (Geyer *et al*, 1990). *Atf-7*^{-/-} mice displayed lower levels of PPI of the acoustic startle response (Figure 1E and Supplementary Figure S2D). Although the association between the startle response and PPI is not currently clear, a negative correlation between the startle response and PPI in WT mice has been

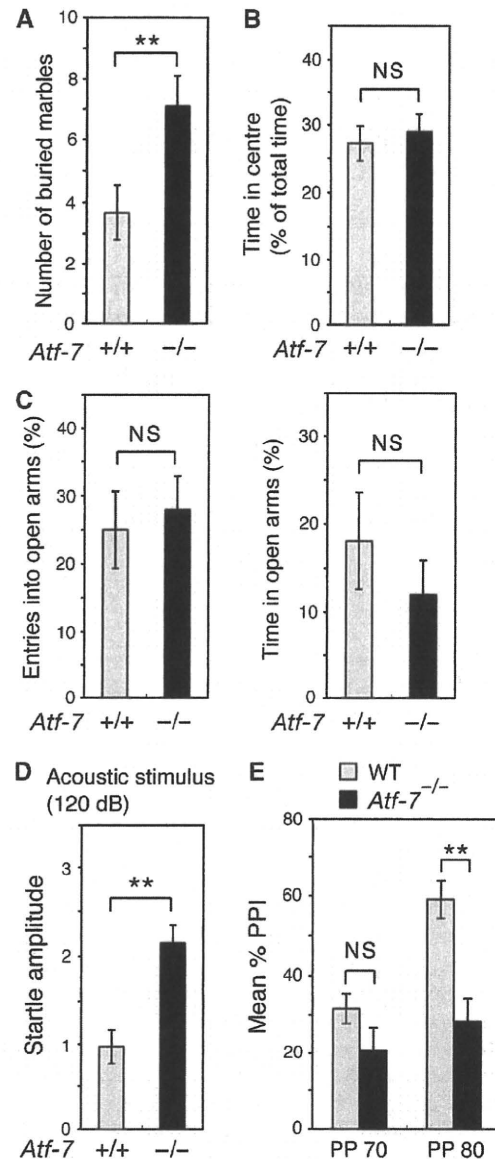


Figure 1 Abnormal behaviours in *Atf-7*^{-/-} mice. Wild-type (WT; +/+) and *Atf-7*^{-/-} C57BL/6 congenic mice were used for all assays. Data are mean ± s.e.m. (A) Marble-burying test. ***P* < 0.01 (*n* = 10–12 for each group). (B) Center of the open-field test. Time spent in the center of the test apparatus is expressed as a percent of total time (10 min). NS, no significant difference (*n* = 13–16 for each group). (C) Elevated plus-maze test. Mice were observed in an elevated plus-maze for 5 min. Percentage of entries into open arms (left) and the time spent in open arms (right) are shown (*n* = 13–16 for each group). (D) Acoustic startle response. Amplitude of the startle response to a 120 dB acoustic stimulus is shown (*n* = 13–16 for each group). (E) Pre-pulse inhibition of the acoustic startle response. The response to a white noise stimulus of 120 dB after a 20 ms pre-pulse warning stimulus (70 or 80 dB) is shown (*n* = 13–16 for each group).

reported (Egashira *et al*, 2005). If these two phenomena are correlated in *Atf-7*^{-/-} mice, an increase in startle reactivity may lead to decreased PPI. However, we cannot exclude the possibility that ATF-7 is independently involved in the modulation of startle response and its PPI.

The *Atf-7*^{-/-} and WT mice responses were indistinguishable in other behavioural tests. We examined spontaneous locomotor activity in a new environment by placing mice in an open-field chamber and monitoring their behaviour. There was no significant difference in the locomotor activity of *Atf-7*^{-/-} and WT mice on the first and second day of the trials (Supplementary Figure S3A and B). We also examined motor coordination using a rotating rod treadmill. Overall, the amount of time mice spent on the rotarod increased with training (Supplementary Figure S3C). The retention time of *Atf-7*^{-/-} mice on the rod was not significantly different from that of WT mice at 0 (stationary), 5, or 10 r.p.m. In the footprint test, there was no significant difference in the stride length and the step width between mutant and WT mice (Supplementary Figure S3D).

In the forced swimming test, there was also no difference between *Atf-7*^{-/-} and WT mice (Supplementary Figure S4A). We also examined spatial learning ability using the Morris Water Maze task. WT and *Atf-7*^{-/-} mice took similar lengths of time to reach the visual platform to escape from the water (Supplementary Figure S4B), thus indicating that *Atf-7*^{-/-} mice have normal vision, motor function, and escape behaviour in the water maze task. Mice were then trained in a hidden platform task, in which mice search for a submerged platform to escape from the water. *Atf-7*^{-/-} and WT mice took similar lengths of time over the 7 days of testing to locate the hidden platform (Supplementary Figure S4C). Thus, *Atf-7*^{-/-} and WT mice were able to learn the location of a hidden platform during the course of the trials. We then carried out a probe test, in which the platform is removed from the pool after completion of the hidden platform task, and the trained mice are allowed to swim freely for 60 s. The time spent in the target quadrant by *Atf-7*^{-/-} mice was similar to that of WT mice (Supplementary Figure S4D). These results indicate that a normal spatial learning ability is present in *Atf-7*^{-/-} mice. The number of crossings of the hidden platform and the swimming distance of *Atf-7*^{-/-} and WT mice was also similar during the probe trial (Supplementary Figure S4E and F). Thus, the performance of *Atf-7*^{-/-} mice in the hidden platform task was indistinguishable from WT mice.

Upregulation of the *Htr5b* gene in the dorsal raphe nucleus of *Atf-7*^{-/-} mice

Atf-7 mRNA was detected in the cortex, the cerebellum, the hippocampus, and the brainstem, including the medulla, the pons, and the midbrain of WT mice (Figure 2A), but not of *Atf-7*^{-/-} mice (Supplementary Figure S5). No obvious morphological abnormalities were found in these tissues in *Atf-7*^{-/-} mice (data not shown). Western blotting indicated that ATF-7 expression levels varied in these tissues (Figure 2B). Several studies have linked abnormal marble-burying behaviour to 5-HT function (Jenck *et al*, 1998). Further, disruptions in PPI of the startle response are correlated not only with D2 dopamine and N-methyl-D-aspartate signalling systems, but also with 5-HT (Geyer *et al*, 2001). Therefore, we focused our attention on the dorsal raphe nuclei of the brainstem, where much of the 5-HT in the brain is localized and relatively high levels of ATF-7 are expressed.

To identify the ATF-7 target genes in the brainstem that may have a role in the abnormal behaviour in *Atf-7*^{-/-} mice, we performed a DNA microarray analysis using RNA from the

brainstem of *Atf-7*^{-/-} and WT mice. The results indicate that 25 genes were upregulated and 38 genes downregulated by more than two-fold by the loss of *Atf-7*. Of these ATF-7 target genes, the functions of 11 of the upregulated genes are known, whereas the functions of only 7 of the downregulated genes have been reported. Among these genes, only the *Htr5b* and the ciliary neurotrophic factor receptor (*Cntfr*) genes have been associated with neuronal function. As the 5-HT system appeared to be associated with the abnormal behaviour of *Atf-7*^{-/-} mice as described above, upregulation of the *Htr5b* gene may be associated with the phenotype of *Atf-7*^{-/-} mice. CNTF is a cytokine that has neurotrophic and differentiating effects on cells in the central nervous system, and the CNTF-CNTF receptor system affects motor neurons (Vergara and Ramirez, 2004). However, there has been no report demonstrating a connection between the CNTF system and anxiety-related behaviours. Therefore, we focused our attention on the *Htr5b* gene.

As *Htr5b* is thought to act as an autoreceptor (Serrats *et al*, 2004), its upregulation may lead to a decrease in the extracellular concentration of serotonin (5-HT). There is abundant evidence for the role of decreased 5-HT in depression and anxiety disorders (Artigas *et al*, 1996). Selective serotonin re-uptake inhibitors (SSRIs), which increase the extracellular concentration of 5-HT in the dorsal raphe nuclei, are widely used as anti-depressant drugs. *Htr5b* mRNA levels in the *Atf-7*^{-/-} brainstem were approximately 12-fold higher than in WT (Figure 2C). *In situ* hybridization showed higher levels of *Htr5b* mRNA expression in the *Atf-7*^{-/-} dorsal raphe nuclei, which also expressed a serotonin transporter mRNA, than in the WT (Figure 2D and Supplementary Figure S6). There appeared to be no obvious difference in the levels of *Htr5b* mRNA between WT and *Atf-7*^{-/-} mice in other regions, including the hippocampus, the habenular nucleus, and the inferior olivary nucleus (Supplementary Figure S7).

Among the many reported antagonists of 5-HT receptors, methiothepin has a relatively high affinity for the 5-HT 5B receptor, although it also binds to other 5HT receptors, including the 5-HT 1A receptor (Boess and Martin, 1994). Injection of methiothepin into *Atf-7*^{-/-} mice suppressed the increased marble-burying behaviour, whereas saline-treated control *Atf-7*^{-/-} mice still exhibited increased marble-burying behaviour (Figure 2E). Furthermore, methiothepin also alleviated the lower levels of PPI in *Atf-7*^{-/-} mice (Figure 2F). These results suggest that increased expression of *Htr5b* mRNA in the *Atf-7*^{-/-} brainstem may, at least partly, contribute to their abnormal behaviour, although loss of ATF-7 could cause changes in the expression of other genes in various regions of the brain and also contribute to abnormal behaviours.

Silencing of the *Htr5b* gene by ATF-7 via direct binding to its 5'-region

Analysis of the DNA sequence in the 5' region of the mouse *Htr5b* gene identified three CRE-like sites at nucleotides -3374, -3340, and -2325 (where +1 is the major transcriptional start site), all of which have only a 1 or 2 bp difference from the consensus CRE sequence (Figure 3A). Gel mobility-shift assays were carried out using nine DNA probes, which cover approximately 4.4 kbp of the 5'-region of *Htr5b*, and nuclear extracts from 293T cells that were transfected with a Flag-tagged ATF-7 expression plasmid or control

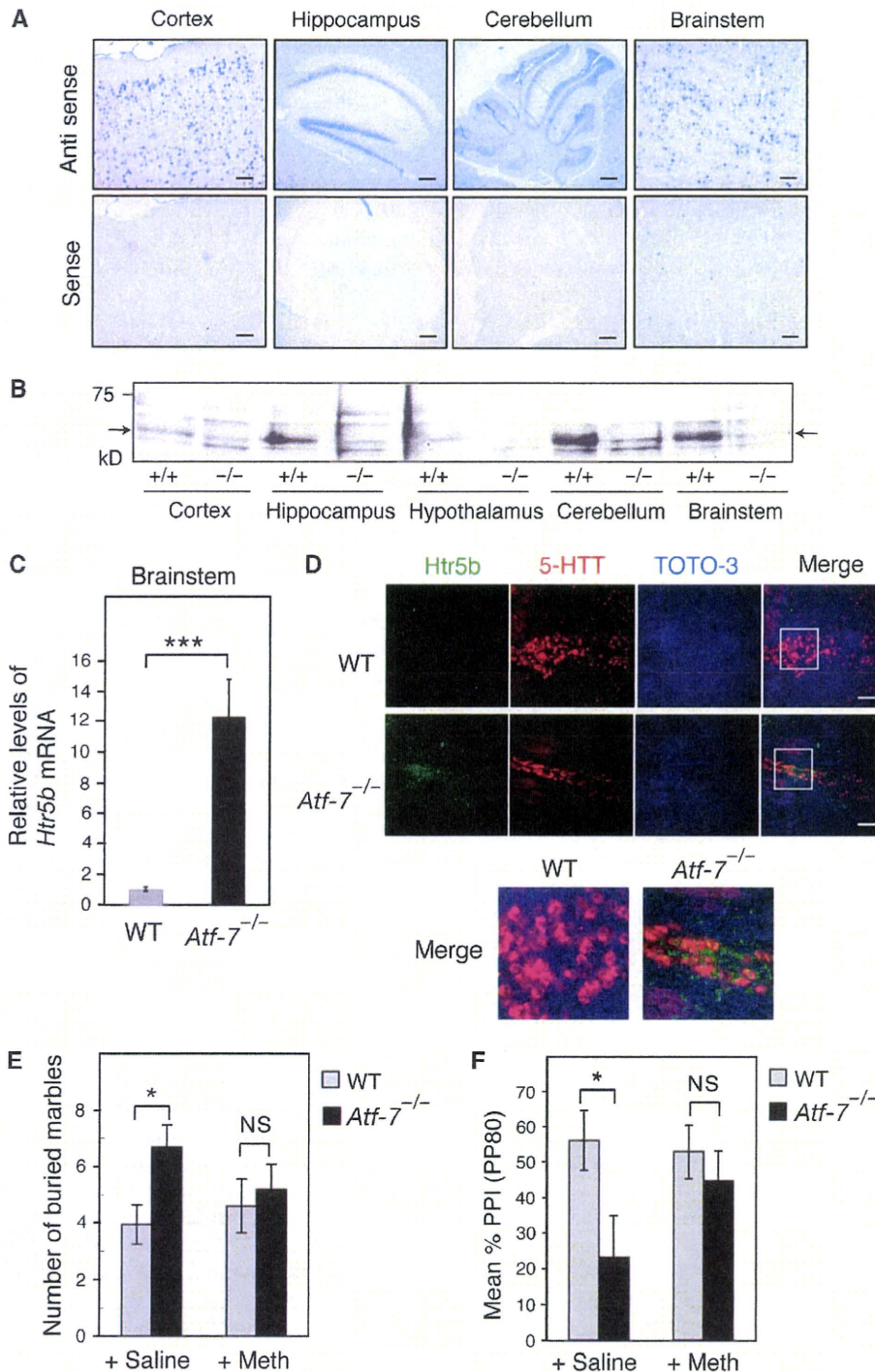


Figure 2 Increased levels of 5-HT receptor 5B (*Htr5b*) mRNA in the *Atf-7*^{-/-} brainstem. (A) *Atf-7* mRNA expression in various regions of the brain was examined by *in situ* hybridization with anti-sense and sense probes. Bar, 100 μ m. (B) Extracts (20 μ g of protein) from the indicated regions of wild-type (WT) or *Atf-7*^{-/-} brains were used for western blotting with anti-ATF-7. The bands indicated by arrows are the ATF-7 signals. (C) Real-time RT-PCR analysis of *Htr5b* mRNA levels using total RNA from the brainstem ($n=4$). *** $P<0.001$. (D) *Htr5b* mRNA expression in the dorsal raphe nuclei was examined by *in situ* hybridization using probes for *Htr5b* (green) and the serotonin transporter (5-HTT, red). Cell nuclei were identified by DNA staining using TOTO-3 (blue). The sections were examined by laser confocal microscopy, and representative images are presented. The panels at the right show the merged images. Bar, 100 μ m. The white box indicates a subregion of each image that is presented at higher magnification below. (E, F) A 5-HT 5B receptor antagonist reduced the abnormal behaviour of *Atf-7*^{-/-} mice. Marble-burying behaviour (E) and pre-pulse inhibition (PPI) (F) of WT and *Atf-7*^{-/-} C57BL/6 congenic mice was examined after administration of either vehicle or methiothepin ($n=10-12$ for each group in E, and $n=7$ for each group in F). * $P<0.05$.

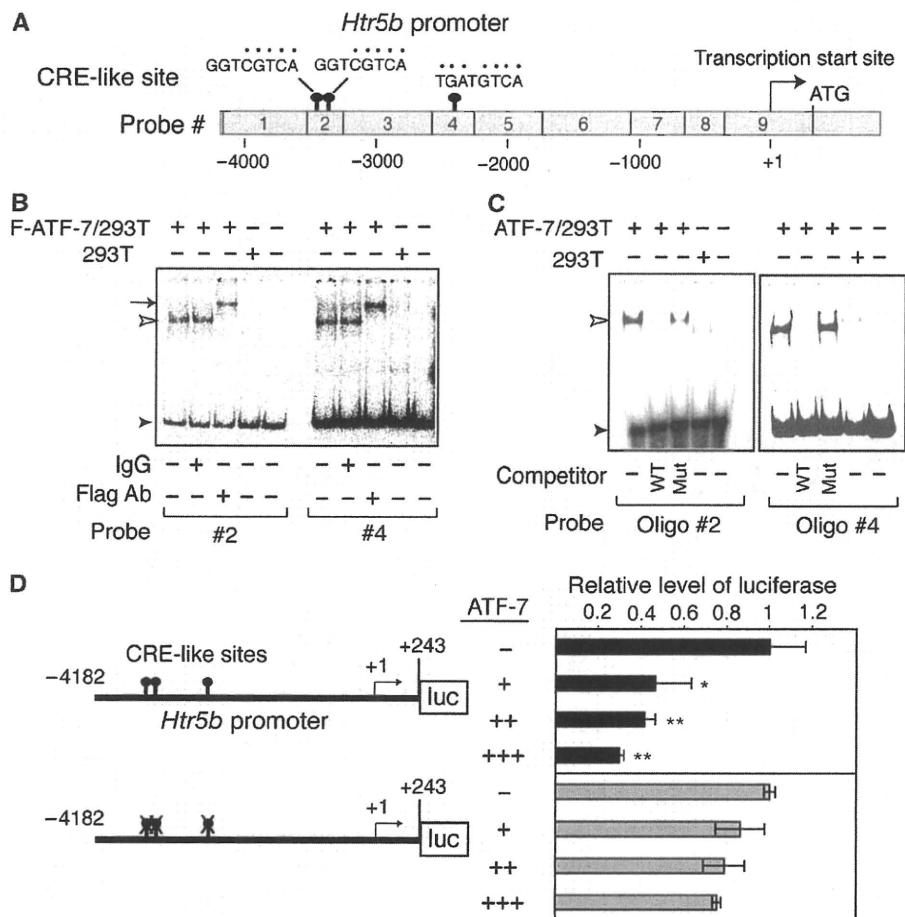


Figure 3 Binding of ATF-7 to the 5-HT receptor 5B (*Htr5b*) promoter region leads to silencing. (A) Presence of cAMP response element (CRE)-like sites in the 5' region of the mouse *Htr5b* gene. The CRE-like sites in the 5'-region of mouse *Htr5b*, and nine DNA probes used for gel mobility-shift assays are shown. (B, C) Gel mobility-shift assays were performed using nuclear extracts prepared from 293T cells transfected with a Flag-ATF-7 expression vector or control empty vector. The #2 and #4 DNA probes were used as the probes in (B). In some lanes, anti-Flag or control IgG was added. In (C), oligonucleotides containing the two CRE-like sites derived from probe #2 or the one CRE-like site from probe #4 were used as probes. In some lanes, a 50-fold excess of competitor containing the same sequence as the probe (wild-type (WT)), or a mutated CRE-like site, was added. Free probe is indicated by a closed arrowhead, whereas ATF-7-bound DNA is shown by an open arrowhead. The ATF-7-DNA complex, which was super-shifted by the anti-Flag antibody, is indicated by the arrow. (D) ATF-7 represses *Htr5b* gene transcription. RN46A cells were transfected with the indicated *Htr5b* promoter-luciferase construct together with 1 (+), 2 (+ +), or 3 (+ + +) μ g of the ATF-7 expression plasmid, or the control empty vector (-), and luciferase activity was measured. Values indicate mean \pm s.d. ($n=3$). * $P<0.05$, ** $P<0.01$.

empty vector. When the #2 or #4 probes were used, a retarded band was detected in extracts containing Flag-ATF-7 (Figure 3B). These specific retarded bands were further shifted when an anti-Flag antibody was added, indicating that the bands contained Flag-ATF-7. In contrast, no retarded bands were observed with the other probes (Supplementary Figure S8). We then used 54 bp and 18 bp oligonucleotides containing the CRE-like sites derived from the #2 and #4 probes, respectively. The retarded bands generated using either probe were competed out by excess amounts of unlabelled competitor oligonucleotide, but not by competitors, which contained mutated CRE-like sites (Figure 3C). These results indicate that ATF-7 binds directly to the CRE-like sites in the 5'-region of the mouse *Htr5b* gene.

When a *Htr5b* promoter-luciferase reporter containing the 4.4 kb 5'-region of the *Htr5b* gene was cotransfected into RN46A cells, which are derived from rat medullary raphe nucleus cells, ATF-7 inhibited luciferase expression by ap-

proximately 70% (Figure 3D). In contrast, mutation of the three CRE-like sites in this reporter relieved the ATF-7-dependent silencing. These results suggest that ATF-7 suppresses the transcription of *Htr5b* through interaction with CRE-like sites.

The results of chromatin immunoprecipitation (ChIP) assays using the brainstem chromatin and an anti-ATF-7 antibody indicated that ATF-7 bound to the region containing CRE-like sites of the *Htr5b* gene, but not to the RNA start site or the 2nd exon (Figure 4A). Further, the binding of ATF-7 to this region was not detected using the *Atf-7*^{-/-} brainstem chromatin. Binding of ATF-2 to this region was also not detected (Figure 4B).

ATF-7 mediates histone H3-K9 trimethylation of the *Htr5b* promoter region by recruiting the ESET HMTase

The results of ChIP assays using the WT brainstem chromatin and anti-H3-K9m3 antibodies indicated that histone H3 in the

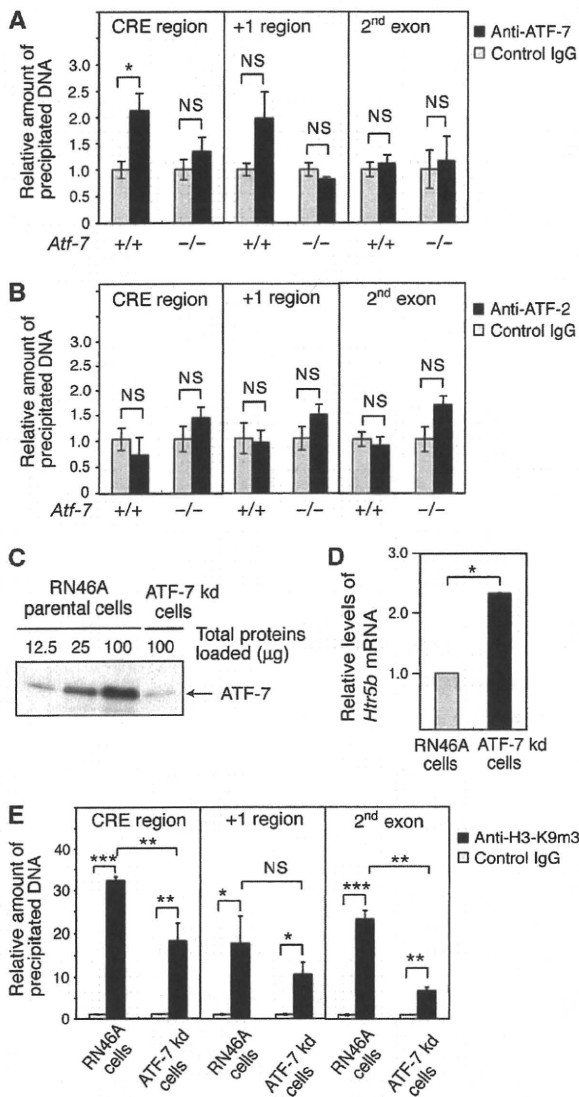


Figure 4 Binding of ATF-7 to the 5-HT receptor 5B (*Htr5b*) promoter is correlated with histone H3-K9 trimethylation. (A, B) Chromatin immunoprecipitation (ChIP) assays were carried out using the brainstem of wild-type (WT) and *Atf-7*^{-/-} mice, and anti-ATF-7 (A), anti-ATF-2 (B), or control IgG (A, B). Extracted DNA was amplified by real-time PCR using primers that cover the cAMP response element (CRE)-like sites, the transcription start site, or the 2nd exon of the *Htr5b* gene. The relative densities of bands are indicated, and each bar represents the mean \pm s.d. ($n = 3$). (C) Generation of an RN46A cell line in which ATF-7 levels are downregulated (ATF-7 kd-RN46A) by expression of a small hairpin-type double-stranded RNA. Nuclear extracts of the parental RN46A cells and the ATF-7 kd-RN46A cells were used for western blotting to detect ATF-7. (D) Real-time RT-PCR analysis of *Htr5b* mRNA levels using RNAs from the parental RN46A cells and ATF-7 kd-RN46A cells. Values are mean \pm s.d. ($n = 3$). (E) ChIP assays were carried out using anti-histone H3-K9m3 and parental RN46A cells or ATF-7 kd-RN46A cells. Extracted DNA was amplified by real-time PCR using primers that cover the CRE-like sites, the transcription start site, or the 2nd exon of the *Htr5b* gene. The relative densities of the bands are indicated, and each bar represents the mean \pm s.d. ($n = 3$). * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$.

region containing the CRE-like sites, the RNA start site, and the 2nd exon of *Htr5b* is trimethylated at K9 (Supplementary Figure S9A). When the *Atf-7*^{-/-} brainstem chromatin was used, similar levels of histone H3-K9 trimethylation were

detected. This result may indicate that *Htr5b* gene transcription is repressed by histone methylation in most cells of the brainstem with the exception of the dorsal raphe nucleus in *Atf-7*^{-/-} mice. To assess the role of ATF-7 in histone H3-K9 methylation, we generated an ATF-7 knock-down (kd) RN46A cell line by expressing a small hairpin-type double-strand RNA. The ATF-7 level was approximately one-eighth that of the parental cell line (Figure 4C). In kd-RN46A cells, *Htr5b* mRNA levels increased approximately 2.3-fold compared with the parental cell line (Figure 4D). In ChIP assays, binding of ATF-7 to the region containing the CRE-like sites of the *Htr5b* gene was detected in parental RN46A cells, but not in the ATF-7 kd-RN46A cells (Supplementary Figure S9B). Binding of ATF-2 to the same region was not detected (Supplementary Figure S9C). The degree of histone H3-K9 trimethylation in the region containing the CRE-like sites or the 2nd exon of *Htr5b* gene was lower in the kd-RN46A cells than in parental cells (Figure 4E). Thus, ATF-7 contributes to H3-K9 trimethylation at the *Htr5b* gene promoter.

To investigate whether the ESET HMTase is involved in the silencing of *Htr5b* by ATF-7, we examined ATF-7-ESET interactions by co-immunoprecipitation. Immunocomplexes prepared from RN46A cell lysates using anti-ATF-7 contained ESET, whereas immunocomplexes prepared with control IgG did not contain ESET (Figure 5A). The results of ChIP assays using the WT brainstem chromatin and anti-ESET antibodies indicated that ESET bound to the region containing CRE-like sites of *Htr5b*, but not to the region containing the RNA start site or the 2nd exon (Figure 5B). Further, binding of ESET to this region was not detected using the *Atf-7*^{-/-} brainstem chromatin. Similar results were also obtained in ChIP assays using the parental RN46A and ATF-7 kd-RN46A cells (Figure 5C).

Members of the ATF-2 subfamily are activated in response to various stresses, and TNF- α is one of the typical inflammatory cytokines which can activate ATF-2 (Brinkman *et al*, 1999). When RN46A cells were treated with TNF- α , the level of *Htr5b* mRNA was gradually increased, by approximately 2.5-fold at 8 h after TNF- α addition (Figure 5D). Phosphorylation of ATF-7 at Thr-53 and of ATF-2 at Thr-71 was also enhanced by TNF- α treatment (Figure 5E). The gradual increase in ATF-7 and ATF-2 phosphorylation observed here is apparently different from the rapid and transient induction of ATF-2 phosphorylation in response to osmotic stress and TGF- β treatment in non-neuronal cells, in which ATF-2 phosphorylation peaks at 30 min after treatment and then decreases (Sano *et al*, 1999). TNF- α also rapidly induces phosphorylation of ATF-2, within 1 h in non-neuronal cells (Brinkman *et al*, 1999). Thus, the gradual response of the p38-ATF-2/7 pathway to TNF- α may be characteristic of neurons. Furthermore, the results of ChIP assays using TNF- α -treated RN46A cells indicate that ATF-7 was released from the 5'-region of *Htr5b* by TNF- α treatment (Figure 5F).

Social isolation stress induces abnormal behaviours similar to those of *Atf-7*^{-/-} mice and *Htr5b* expression

Defects in PPI attenuation of the acoustic startle response are induced by social isolation stress (Wilkinson *et al*, 1994), although isolated animals exhibit a variety of phenotypes, including aggressive behaviour (Blanchard *et al*, 2001). This observation suggests that exposure of WT mice to social

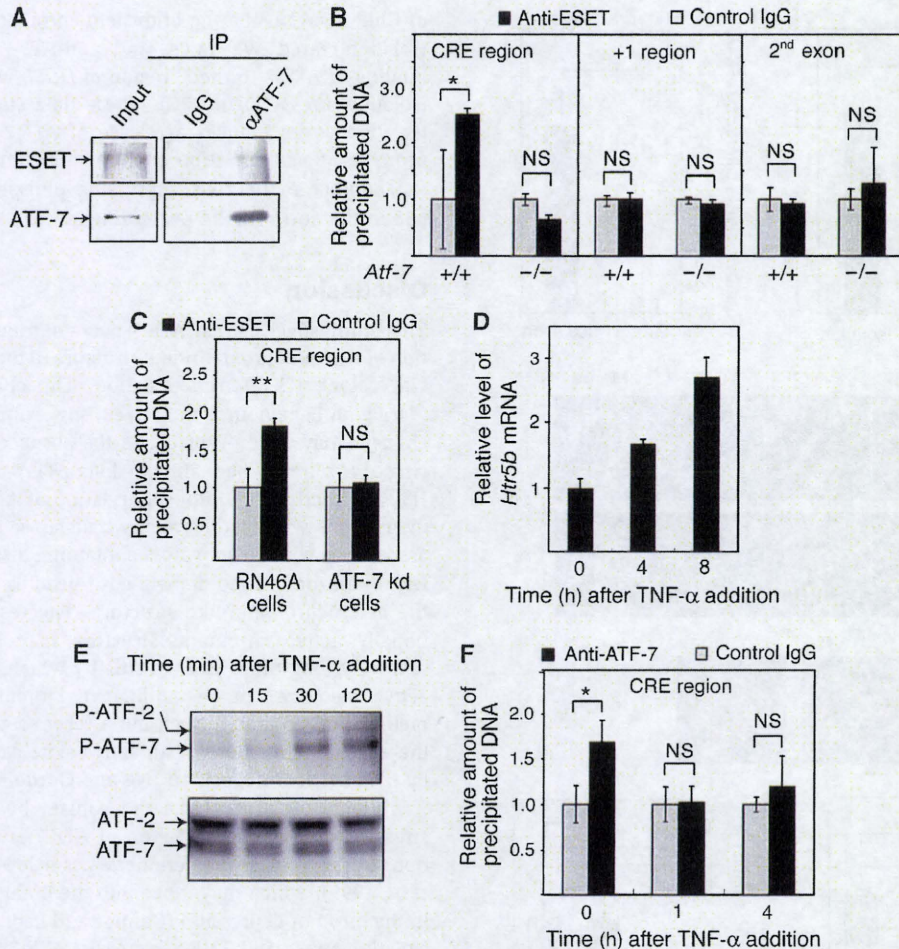


Figure 5 ATF-7 recruits the ESET HMTase to the 5-HT receptor 5B (*Htr5b*) gene. (A) Co-immunoprecipitation of ATF-7 and ESET. Whole cell lysates of RN46A cells were immunoprecipitated with anti-ATF-7 or control IgG, and the immunocomplexes were then analyzed by western blotting using anti-ESET or anti-ATF-7 antibodies. (B) Recruitment of ESET to the *Htr5b* gene by ATF-7. Chromatin immunoprecipitation (ChIP) assays were carried out using anti-ESET and chromatin from wild-type (WT) or *Atf-7*^{-/-} brainstems. Extracted DNA was amplified by real-time PCR using primers that cover the cAMP response element (CRE)-like sites, the RNA start site, or the 2nd exon of the *Htr5b* gene. The relative densities of the bands are indicated, and each bar represents the mean \pm s.d. ($n=3$). (C) ChIP assays were carried out using anti-ESET and the parental RN46A or the ATF-7 kd-RN46A cells. Extracted DNA was amplified by real-time PCR using primers that cover the ATF-7-binding sites of the *Htr5b* gene ($n=3$). (D) Upregulation of *Htr5b* mRNA by TNF- α in RN46A cells. Real-time RT-PCR analysis of *Htr5b* mRNA was performed using RNAs from RN46A cells treated with TNF- α for the indicated times ($n=3$). (E) Increase in phosphorylation of ATF-7 in response to TNF- α . Nuclear extracts were prepared from RN46A cells treated with TNF- α for the indicated times, and used for western blotting with antibodies that recognize the indicated proteins. We have identified the ATF-7 band by confirming that it was lost in the *Atf-7*^{-/-} whole brain nuclear extract (Supplementary Figure S10). (F) Release of ATF-7 from the *Htr5b* promoter by TNF- α treatment. ChIP assays were carried out as described above using anti-ATF-7 and RN46A cells treated with TNF- α for the indicated times ($n=3$). * $P<0.05$, ** $P<0.01$.

isolation stress may cause abnormal behaviour and an increase in *Htr5b* mRNA levels in the dorsal raphe nuclei, both of which were observed in *Atf-7*^{-/-} mice. In fact, WT mice exhibited increased marble-burying behaviour after 1 month of isolation rearing (Figure 6A). Furthermore, after isolation rearing of WT mice for 1 month, *Htr5b* mRNA levels in the brainstem increased approximately 12-fold (Figure 6B). *In situ* hybridization indicated that the level of *Htr5b* mRNA in the dorsal raphe nuclei, which also expressed serotonin transporter mRNA, was enhanced by isolation rearing (Figure 6C and Supplementary Figure S6C). The result of ChIP assays using the brainstem from group- or isolation-reared mice indicated that the degree of histone H3-K9 trimethylation was not affected by isolation stress (Supplementary Figure S11). This result may indicate that *Htr5b* gene transcription is repressed by histone methylation

in most brainstem cells with the exception of the dorsal raphe nucleus in isolation-reared mice.

When *Atf-7*^{-/-} mice were exposed to isolation stress, enhancement of marble-burying behaviour was observed (Figure 6D). However, *Htr5b* mRNA levels in the *Atf-7*^{-/-} brainstem were not further increased by isolation stress (Figure 6E). These results suggest that isolation stress induces abnormal marble-burying behaviour not only by induction of *Htr5b* but also by changing the expression of other genes.

Social isolation stress induces the phosphorylation of ATF-7 and release of ATF-7 from the *Htr5b* gene

We examined phosphorylated ATF-7 (P-ATF-7) signals in the dorsal raphe nucleus of group- and isolation-reared WT mice. Social isolation stress significantly increased the number of

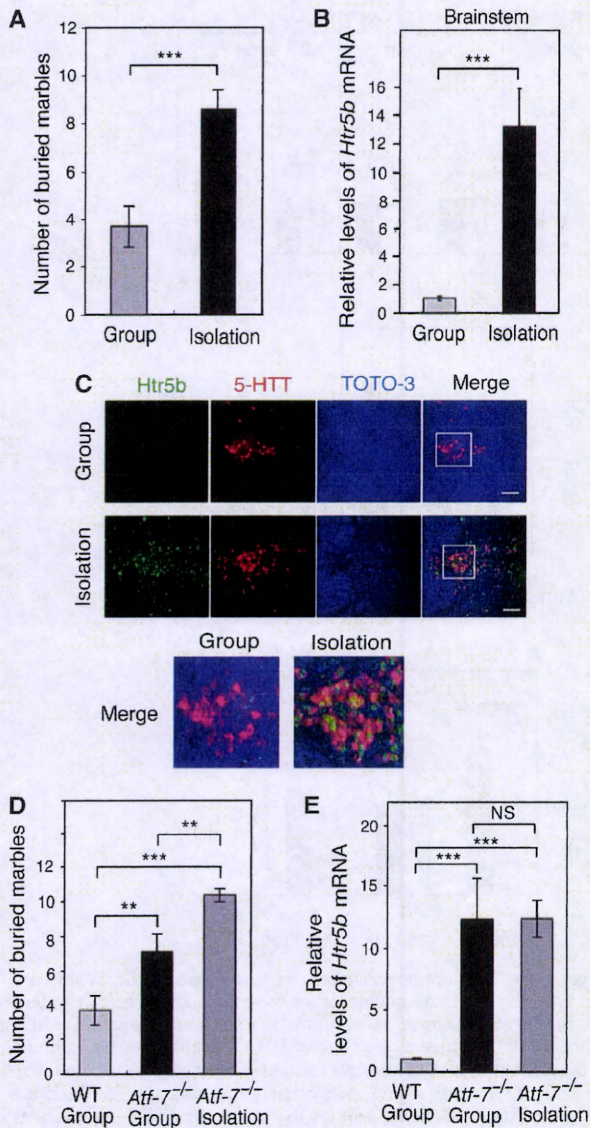


Figure 6 Social isolation stress increases marble-burying behaviour and induces 5-HT receptor 5B (*Htr5b*) mRNA expression. (A) Marble-burying behaviour of group- and isolation-reared WT mice was examined ($n = 10$ – 12 of C57BL/6 congenic mice for each group). (B) Real-time RT-PCR analysis of *Htr5b* mRNA expression was performed using RNAs from the brainstem of group- and isolation-reared WT mice ($n = 3$). (C) *Htr5b* mRNA expression in the dorsal raphe nuclei of group- and isolation-reared WT mice was examined by *in situ* hybridization, as described in Figure 2D. (D) The group-reared WT and *Atf-7*^{-/-} mice, and the isolation-reared *Atf-7*^{-/-} mice were used for marble-burying tests ($n = 10$ – 12 of C57BL/6 congenic mice for each group). (E) Real-time RT-PCR analysis of *Htr5b* mRNA was performed using RNAs from the brainstems of the mice described in D ($n = 3$). ** $P < 0.01$, *** $P < 0.001$.

neurons expressing P-ATF-7 and P-ATF-2 in the dorsal raphe nucleus (Figure 7A). As the amino-acid sequence of the p38 phosphorylation site is highly conserved between ATF-7 and ATF-2, antibodies to P-ATF-7 also recognize P-ATF-2. However, the results of western blotting using extracts from the brainstem indicated that the phosphorylation of ATF-7 increased under social isolation stress conditions, whereas the phosphorylation of ATF-2 did not (Figure 7B). The results

of ChIP assays using the brainstem chromatin of group- and isolation-reared WT mice, and anti-ATF-7, indicated that binding of ATF-7 to the 5'-region of *Htr5b* was lost by social isolation stress (Figure 7C). These data suggest that social isolation stress induces a release of ATF-7 from the *Htr5b* promoter. Social isolation stress also significantly increased the number of neurons expressing phosphorylated, active p38 in the dorsal raphe nuclei (Figure 7D).

Discussion

This study suggests that ATF-7 may contribute to the formation of a heterochromatin-like structure in the *Htr5b* promoter via histone H3-K9 trimethylation. The effect of isolation-rearing on behaviour is long-lived, suggesting that epigenetic changes may have occurred. In the absence of stress, silencing of *Htr5b* may be maintained via ATF-7-mediated histone H3-K9 trimethylation. Phosphorylation of ATF-7 may disrupt interactions with histone methyltransferase and enhance the association with co-activators containing histone acetyltransferase and/or histone demethylase, leading to disruption of the heterochromatin-like structure. The resulting transcriptionally active chromatin structure may be stable for a relatively long time. Mechanisms by which transcriptionally active memory can be modulated without affecting DNA methylation remain elusive, but a recent report showed that the epigenetic memory of an active state can be established by histone H3.3 deposition (Ng and Gurdon, 2008).

Although at present the mechanism by which isolation stress induces phosphorylation of ATF-7 is unknown, isolation stress increases the peripheral tissue levels of TNF- α (Wu *et al*, 1999), which may move into the brain and be involved in the onset of depression (Connor and Leonard, 1998). Both isolation stress and TNF- α increased ATF-7 phosphorylation and *Htr5b* mRNA levels, which are accompanied by a release of ATF-7 from the *Htr5b* promoter. The mechanism by which the phosphorylation of ATF-7 causes a release of ATF-7 from target sites is unknown. The phosphorylation-induced release of ATF-7 from DNA could be caused by changes in interactions between ATF-7 and uncharacterized factors that enhance ATF-7 affinity for DNA. ATF-7 is highly homologous to ATF-2, but there are differences between the two proteins. In luciferase reporter assays, ATF-2 activated transcription from the CRE-containing promoter, but ATF-7 did not, even in the presence of active p38. This result suggests that ATF-7 may have a role primarily in transcriptional repression. As we have observed that ATF-7 forms a heterodimer with ATF-2 (data not shown), further study is required to compare the function of ATF-7 and ATF-2 homodimers and ATF-7-ATF-2 heterodimers.

Our results suggest that upregulation of *Htr5b* may partly contribute to the abnormal behaviour of *Atf-7*^{-/-} mice. The rodent 5-HT5 receptor family consists of two receptors, 5-HT 5A and 5B (*Htr5a* and *Htr5b*), which share 69% amino acid identity and have 23–34% homology with other 5-HT receptors (Plassat *et al*, 1992). *Htr5a* has been identified in mouse, rat, and human. Mouse and rat have a functional *Htr5b* gene, whereas the human coding sequence is interrupted by stop codons (Grailhe *et al*, 2001). Thus, humans do not have a functional *Htr5b*, and, therefore, another subtype, such as *Htr5a*, could be upregulated and has a role in response to social isolation stress in humans. *Htr5a*-deficient mice

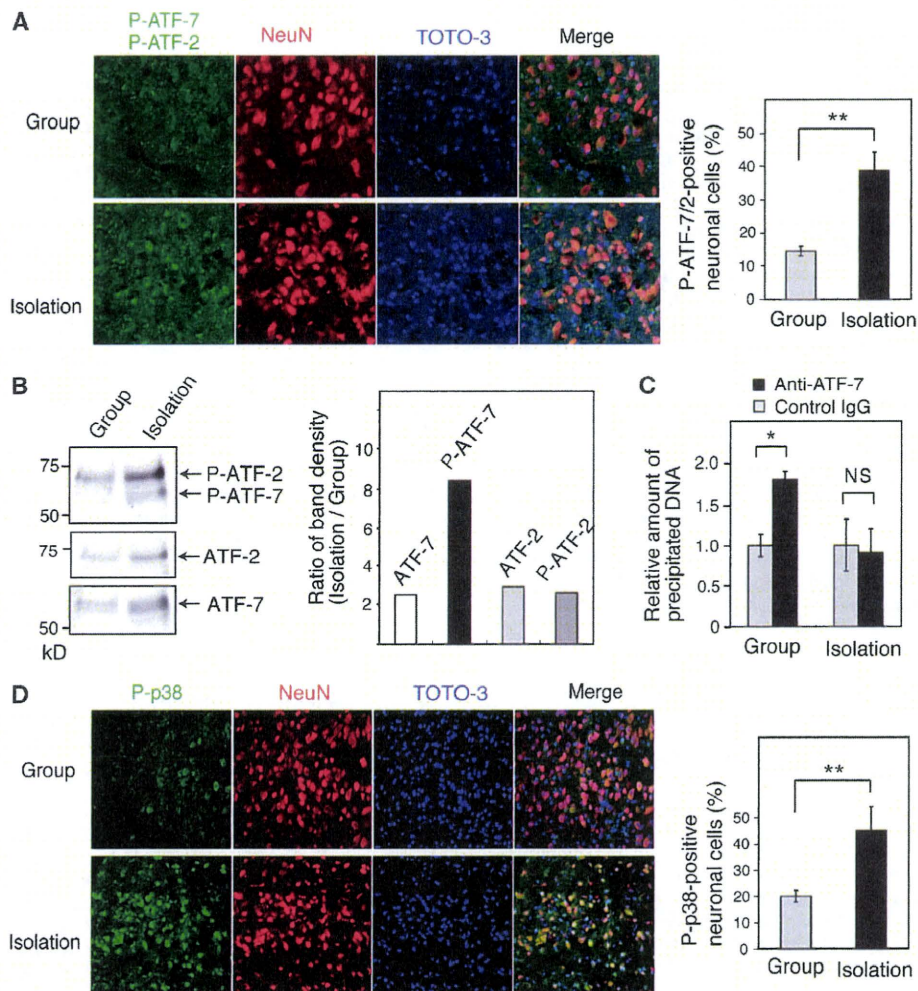


Figure 7 Social isolation stress induces ATF-7 phosphorylation and release of ATF-7 from the 5-HT receptor 5B (*Htr5b*) promoter. (A) Brain sections containing dorsal raphe nuclei of group- or isolation-reared wild-type (WT) mice were stained with antibodies which recognize P-ATF-7 and P-ATF-2 (green), or NeuN (red), a neuronal specific nuclear protein. DNA was stained with TOTO-3 (blue). The merged images are shown in the right panels. The average number of neurons expressing P-ATF-7 or P-ATF-2 in three independent experiments is indicated by the bar graph \pm s.d. (B) Nuclear extracts were prepared from the brainstem of group- or isolation-reared WT mice, which were perfused with PFA, and analyzed by SDS-PAGE after decrosslinking, followed by western blotting with anti-P-ATF-2/7, anti-ATF-7, and anti-ATF-2. The ratio of the density of each band in isolation-reared mice to that in group-reared mice is indicated in the bar graph. (C) Release of ATF-7 from the *Htr5b* promoter by isolation stress. Chromatin immunoprecipitation (ChIP) assays were carried out using chromatin from the brainstem of group- or isolation-reared WT mice, and anti-ATF-7. Extracted DNA was amplified by real-time PCR using primers that cover the ATF-7-binding sites of *Htr5b* ($n = 3$). (D) Brain sections containing dorsal raphe nuclei of group- or isolation-reared WT mice were stained with antibodies that recognize P-p38, as described above. The number of neurons expressing P-p38 is quantified at the right ($n = 3$). * $P < 0.05$, ** $P < 0.01$.

display increased exploratory activity when exposed to new environments, suggesting that *Htr5a* modulates the activity of neural circuits involved specifically in exploratory behaviour (Grailhe *et al*, 1999). However, there has been no report of a mouse knockout of *Htr5b*.

Isolation-rearing was recently reported to reduce (27.0–60.9%) transcription of many postsynaptic 5-HT receptors in the prefrontal cortex and the 5-HT 1B, 2A, and 2C receptors in the hypothalamus and the midbrain, whereas 5-HT 6 receptor mRNA levels increased (52.5%) in the hippocampus (Bibancos *et al*, 2007). *Htr5b* expression was not examined. The fold change in mRNA level of these 5-HT receptors was much lower than the change in *Htr5b* mRNA described herein (12-fold). We have also examined the expression level of 5-HT receptors other than 5B in *Atf-7*^{-/-}

mice. Slight decreases in the 2A and 2C receptors in the hypothalamus and 1A and 1B receptors in the cortex were observed in *Atf-7*^{-/-} mice, whereas the levels of 2A and 3A receptors in the cortex increased slightly (Supplementary Figure S12). However, the degree of these changes was much lower than the change in *Htr5b* in the brainstem in response to social isolation rearing. There was no difference in the *Htr5b* mRNA levels in the hypothalamus and in the cortex between WT and *Atf-7*^{-/-} mice. Thus, the degree of change in the *Htr5b* mRNA levels in the brainstem in response to loss of ATF-7 or social isolation stress is dramatic, suggesting a unique role for *Htr5b*. However, it is also likely that loss of ATF-7 changes the expression of multiple genes in various regions of the brain that may also contribute to abnormal behaviours.

Materials and methods

Animals

All mice used were 2–6-month-old males. In the original behaviour tests, immunohistochemistry, RNA analysis, and ChIP assays, the littermate mice, which were generated by mating *Atf-7* heterozygotes with a mixed CBA (25%) × C57BL/6 (75%) genetic background, were used. To confirm some behavioural abnormalities of *Atf-7*^{-/-} mice, C57BL/6 congenic mice, which were generated by backcrossing onto a C57BL/6 genetic background for 7 generations, were used. Both WT and *Atf-7*^{-/-} mice were maintained in a temperature and humidity-controlled room with free access to food and water. Animals were maintained on a 12-h-light and 12-h-dark cycle (lights on at 0800 h, lights off at 2000 h). For social isolation stress, mice were group-housed until 1-month old and then housed individually for 1 month before the start of the experiments. Experiments were conducted in accordance with the guidelines of the Animal Care and Use Committee of the RIKEN Institute.

Behavioural analysis

Marble-burying test. The marble-burying behaviour tests were carried out using C57BL/6 congenic mice (Figures 1A and 2E) or mice with a mixed CBA (25%) × C57BL/6 (75%) genetic background (Supplementary Figure S2A) as described (Yamada *et al*, 2002). The mice were placed individually in plastic cages (20 × 14 × 22 cm³) for 30 min (habituation trial) and then returned to their home cages. Twelve clean, coloured glass marbles (10 mm in diameter) were evenly spaced 3–5 cm apart on 5 cm deep sawdust in the habituation cages. Mice were then re-introduced into these cages without food and water (each test mouse was returned to the same cage in which they had been habituated). The results of marble-burying behaviour were expressed as the number of marbles, at least, two-thirds buried within 30 min.

Acoustic startle response. The acoustic startle response was measured using specific startle chambers (O'Hara & Co. in Figure 1D and E, or SR-LAB, San Diego Instruments in Supplementary Figure S2C and D). The startle chamber consisted of a Plexiglas cylinder 3.5 or 3.8 cm in diameter, resting on a sensor block or on a Plexiglas frame in a sound-attenuated, ventilated enclosure. Acoustic bursts were presented through a loudspeaker mounted 25 or 29 cm above the cylinder. A piezoelectric transducer mounted below the sensor-block/frame detected motion of the animal in the cylinder. Stabilimeter readings were rectified and recorded by a microcomputer and interface ensemble (O'Hara & Co. or San Diego Instruments). One mouse was placed in each chamber and allowed to acclimate for 10 min and then the experimental session was started. Background noise was set at 65 dB white noise throughout both the acclimation period and the session. In a session, 10 trials for three types of stimuli each (total of 30 trials) were given in pseudo-random order after three initial startle-stimuli (20-ms burst of 120 dB white noise), which were given to avoid the effect of high responses to initial stimulations in the experiments. One type of trial was a pulse-alone (P alone) trial, which involved a 20-ms burst of 120 dB white noise, and the other two types of trials were pre-pulse and pulse (PP70 & P and PP80 & P) trials, in which a 20-ms burst of 70 or 80 dB white noise, respectively, was followed by a 20-ms burst of 120 dB white noise 100 ms later. The inter-trial intervals averaged 40 s (20–60 s) and were pseudo-randomized. The startle response was measured for 300 ms (Figure 1D and E) or 100 ms (Supplementary Figure S2C and D) from the beginning of pulse presentation, and the largest value was defined as the startle amplitude. The startle amplitudes of the animal in response to repetitions of each trial type were averaged across the session. The experimental schedule was controlled by a micro-computer.

Methiothepin treatment. Methiothepin mesylate salt (Sigma-Aldrich) (0.1 mg/kg weight) and saline (Otsuka) were administered by intra-peritoneal injection. Tests were conducted 1 h after drug administration.

Histological analysis and immunohistochemistry

Tissues were fixed by perfusion with 4% PFA, dehydrated, and embedded in paraffin. Sections (5 µm) were stained with hematoxylin and eosin according to standard procedures. Frozen sections

(10 µm) were used for immunohistochemistry. For indirect immunofluorescent staining, anti-p71-ATF-2 (#922, Cell Signaling), anti-p180/p182-p38 MAPK (#4631, Cell Signaling), anti-NeuN (MAB377, Chemicon), and TOTO-3 (Invitrogen) were used. The frozen sections were washed twice with Tris-buffered saline (144 mM NaCl, 10 mM Tris-HCl, pH 7.6) and incubated overnight at 4°C with primary antibody. Biotin-conjugated anti-rabbit IgG antibody served as the secondary antibody and was incubated at room temperature for 2 h and further incubated with streptavidin Alexa Fluor 488 (Molecular Probes) and Alexa Fluor 546 anti-mouse IgG at room temperature for 2 h.

Luciferase reporter assay

The *Htr5b* promoter-luciferase reporter, in which a 4.4 kb mouse *Htr5b* promoter DNA fragment (from -4182 to +243) was linked to the luciferase gene, was constructed. The mutant construct containing mutated CRE-like sites was constructed by replacing the 42-bp region containing the two CRE sites upstream (-3374 ~ -3333) with the 6-bp sequence (GAGCTC) of a *Sad* linker, whereas the 8-bp sequence of the third CRE site downstream (-2325 ~ -2318) was replaced by the 8-bp sequence (AACGCGTT) of a *MtII* linker. To generate the ATF-7 expression vector, pAct-ATF-7, the human ATF-7 cDNA was inserted into the chicken cytoplasmic β-actin promoter-containing vector. RN46A cells were cultured in Dulbecco's modified Eagle's medium (DMEM) and F-12 HAM (D8062, Sigma-Aldrich) (1:1 mixture) supplemented with 10% FBS at 33°C in 5% CO₂. RN46A cells were transfected using Lipofectamine Plus (Invitrogen) with 0.1 µg of the *Htr5b* promoter-luciferase reporter, various amounts of the ATF-7 expression plasmid or the control empty vector (0–3 µg), and 1 µg of the internal control *pras-β-gal*, in which the β-galactosidase gene was linked to the human *c-Ha-ras* promoter. At 48 h post-transfection, luciferase activity was measured and normalized for transfection efficiency by β-galactosidase activity.

ChIP assays

The brain tissues were prepared essentially as described by Tsankova *et al* (2004). The brainstem was removed by gross dissection, minced into ~0.3 mm pieces, and immediately cross-linked in 1.5% formaldehyde for 15 min at room temperature. After addition of glycine to a final concentration of 0.125 M to quench the crosslinking reaction, the chromatin was solubilized and extracted with lysis buffer, and sheared to 400–600 bp fragments by sonication. ChIP assays were carried out essentially as described by Jin *et al* (2006). Immunoprecipitation was carried out for 4–10 h at 4°C with anti-ATF-7 (2F10 or 1A7), anti-histone H3 K9-m3 (ab8898, Abcam), anti-ESET (Upstate #07-378), or normal mouse or rabbit IgG as negative controls. The immunocomplexes were washed and incubated at 65°C in 100 µl of IP elution buffer (1% SDS, 0.1 M NaHCO₃, 250 mM NaCl, 200 µg/ml proteinase K, 10 mM DTT) to release proteins. The free precipitated DNA was further purified using a QIAquick PCR Purification Kit (Qiagen) and eluted in 30 µl sterile water. Eluted DNA samples were used for real-time PCR (7500 Real Time PCR System, Applied Biosystems). ChIP assays using RN46A cells were carried out essentially as described by Jin *et al* (2006). The primers and TaqMan probes (Qiagen) used for amplification are described in Supplementary Table S1.

Co-immunoprecipitation assay

For co-immunoprecipitation assays of endogenous ATF-7 and ESET, RN46A cells were lysed by mild sonication in NETN buffer (20 mM Tris-HCl, pH 8.0, 1 mM EDTA, 0.5% NP40, 400 mM NaCl). Lysates were immunoprecipitated using anti-ATF-7 (1A7) or control IgG. Immunocomplexes were resolved on 10% SDS polyacrylamide gels, and analyzed by western blotting with anti-ESET (Upstate #07-378).

Supplementary data

Supplementary data are available at *The EMBO Journal* Online (<http://www.embojournal.org>).

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Conflict of interest

The authors declare that they have no conflict of interest.

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