Enhanced sensitivity to ethanol-induced inhibition of LTP in CA1 pyramidal neurons of socially isolated C57BL/6J mice: role of neurosteroids

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INTRODUCTION
Ethanol (EtOH) induced impairment of long-term potentiation (LTP) in the rat hippocampus is prevented by the 5α-reductase inhibitor finasteride, suggesting that this effect of EtOH is dependent on the increased local release of neurosteroids such as 3α,5α-THP that promote GABA-mediated transmission. Given that social isolation (SI) in rodents is associated with altered plasma and brain levels of such neurosteroids as well as with an enhanced neurosteroidogenic action of EtOH, we examined whether the inhibitory effect of EtOH on LTP at CA3–CA1 hippocampal excitatory synapses is altered in C57BL/6J mice subjected to SI for 6 weeks in comparison with group-housed (GH) mice. Extracellular recording of field excitatory postsynaptic potentials (fEPSPs) as well as patch-clamp analysis were performed in hippocampal slices prepared from both SI and GH mice. Consistent with previous observations, recording of fEPSPs revealed that the extent of LTP induced in the CA1 region of SI mice was significantly reduced compared with that in GH animals. EtOH (40 mM) inhibited LTP in slices from SI mice but not in those from GH mice, and this effect of EtOH was abolished by co-application of 1 μM finasteride. Current-clamp analysis of CA1 pyramidal neurons revealed a decrease in action potential (AP) frequency and an increase in the intensity of injected current required to evoke the first AP in SI mice compared with GH mice, indicative of a decrease in neuronal excitability associated with SI. Together, our data suggest that SI results in reduced levels of neuronal excitability and synaptic plasticity in the hippocampus. Furthermore, the increased sensitivity to the neurosteroidogenic effect of EtOH associated with SI likely accounts for the greater inhibitory effect of EtOH on LTP in SI mice. The increase in EtOH sensitivity induced by SI may be important for the changes in the effects of EtOH on anxiety and on learning and memory associated with the prolonged stress attributable to SI.

Keywords: social isolation, ethanol, hippocampus, neurosteroids, neuronal excitability, stress, LTP

Ethanol (EtOH) induced impairment of long-term potentiation (LTP) in the rat hippocampus is prevented by the 5α-reductase inhibitor finasteride, suggesting that this effect of EtOH is dependent on the increased local release of neurosteroids such as 3α,5α-THP that promote GABA-mediated transmission. Given that social isolation (SI) in rodents is associated with altered plasma and brain levels of such neurosteroids as well as with an enhanced neurosteroidogenic action of EtOH, we examined whether the inhibitory effect of EtOH on LTP at CA3–CA1 hippocampal excitatory synapses is altered in C57BL/6J mice subjected to SI for 6 weeks in comparison with group-housed (GH) mice. Extracellular recording of field excitatory postsynaptic potentials (fEPSPs) as well as patch-clamp analysis were performed in hippocampal slices prepared from both SI and GH mice. Consistent with previous observations, recording of fEPSPs revealed that the extent of LTP induced in the CA1 region of SI mice was significantly reduced compared with that in GH animals. EtOH (40 mM) inhibited LTP in slices from SI mice but not in those from GH mice, and this effect of EtOH was abolished by co-application of 1 μM finasteride. Current-clamp analysis of CA1 pyramidal neurons revealed a decrease in action potential (AP) frequency and an increase in the intensity of injected current required to evoke the first AP in SI mice compared with GH mice, indicative of a decrease in neuronal excitability associated with SI. Together, our data suggest that SI results in reduced levels of neuronal excitability and synaptic plasticity in the hippocampus. Furthermore, the increased sensitivity to the neurosteroidogenic effect of EtOH associated with SI likely accounts for the greater inhibitory effect of EtOH on LTP in SI mice. The increase in EtOH sensitivity induced by SI may be important for the changes in the effects of EtOH on anxiety and on learning and memory associated with the prolonged stress attributable to SI.
anxiolytic and sedative effects in vivo (Bitran et al., 1993, 1995; Freeman et al., 1993; Picazo and Fernandez-Guasti, 1995; Kokate et al., 1999; Reddy et al., 2004). Plasma and brain levels of GABAergic neuroactive steroids are affected by acute or chronic stress as well as by the acute administration of EtOH in rodents (Purdy et al., 1991; Concasa et al., 1996; Barbaccia et al., 1999; Morrow et al., 1999; Serra et al., 2006).

The steroidogenic action of EtOH is thought to be mediated by stimulation of the hypothalamic–pituitary–adrenal (HPA) axis and results in an increase in the circulating and brain levels of neuroactive steroids. However, EtOH directly stimulates neurosteroidogenesis in rat brain slices and thus independently of peripheral organs (Sanna et al., 2004; Criswell and Breese, 2005; Izumi et al., 2007). EtOH induces the local release of neurosteroids at the synaptic level as well as positive modulation of GABAARs in CA1 pyramidal neurons, with both effects being prevented by the 5α-reductase inhibitor finasteride, suggesting that local production of neurosteroids such as 3α,5α-THP is necessary for modulation of GABAAR function by EtOH. In addition, this neurosteroidal effect of EtOH was shown to result in inhibition of long-term potentiation (LTP) in the CA1 hippocampal region (Izumi et al., 2007). LTP in the hippocampus is a form of synaptic plasticity that provides a consolidated cellular mechanism of memory formation (Mayford, 2007). Membrane excitability in CA1 pyramidal neurons has been found to be related both to the performance of learning tasks (Moyer et al., 2000; Tombaugh et al., 2005) as well as to successful learning (Moyer et al., 1996; Thompson et al., 1996).

Postweaning social isolation (SI) is a well-characterized paradigm of mild prolonged stress that is associated with marked behavioral, neuroendocrine, and neurochemical changes (Hall et al., 1998; Serra et al., 2000, 2005, 2006; Ferdman et al., 2007) and that is dependent on gender (Pietropaolo et al., 2008). A decrease in the brain concentrations of GABAergic neuroactive steroids such as 3α,5α-THP in animals subjected to SI is accompanied by an increased efficacy of EtOH in the stimulation of steroidogenesis and GABAAR function (Serra et al., 2003, 2006). Moreover, SI induces increased voluntary consumption of EtOH in adult C57BL/6J mice (Sanna et al., 2011).

We have now evaluated the effects of SI on neuronal excitability and on the inhibition by EtOH of LTP induction in the CA1 region of the mouse hippocampus as well as the role of neurosteroids in such effects. The decrease in neurosteroid levels associated with SI was found to result in a reduced level of neuronal excitability and an enhanced inhibitory effect of EtOH on LTP in the CA1 hippocampal region.

**MATERIALS AND METHODS**

**ANIMALS**

C57BL/6J mice (Charles River, Como, Italy) were bred in our animal facility and maintained under an artificial 12-h-light, 12-h-dark cycle (lights on from 08:00 to 20:00 h), a constant temperature of 22 ± 2°C, and a relative humidity of 65%. They had free access to water and standard laboratory food at all time. Animal care and handling throughout the experimental procedures were in accordance with the European Communities Council Directive of November 24, 1986 (86/609/EEC). The experimental protocols were also approved by the Animal Ethics Committee of the University of Cagliari.

**STRESS PARADIGM**

Newborn mouse pups were left undisturbed with their mothers until weaning (21 days). After weaning, male mice were randomly assigned to be housed six per cage (group-housed, GH) or one per cage (SI) for 6 weeks. In a separate set of experiments, 12 SI mice were injected once a day subcutaneously with progesterone (5 mg/kg, dissolved in 20% β-cyclodextrin), and six of which were co-treated with finasteride (25 mg/kg, dissolved in 20% β-cyclodextrin, s.c.) throughout the entire 6-week period of isolation; control SI mice received an equivalent injection of the vehicle solution according to the same schedule.

**PREPARATION OF MOUSE HIPPOCAMPAL SLICES**

Hippocampal slices were prepared from GH and SI mice as previously described (Sanna et al., 2011). In brief, the animals were subjected to deep anesthesia with chloroform and decapitated, and the brain was rapidly removed from the skull. For extracellular recordings, the brain was transferred to a standard artificial cerebrospinal fluid (ACSF) containing (in mM): 126 NaCl, 3 KCl, 2 CaCl2, 1 MgCl2, 26 NaHCO3, 1.25 NaH2PO4, and 10 d-glucose (pH 7.4, set by aeration with 95% O2 and 5% CO2). Alternatively, for patch-clamp experiments, the brain was transferred to a modified ACSF containing (in mM): 220 sucrose, 2 KCl, 0.2 CaCl2, 6 MgSO4, 26 NaHCO3, 1.3 NaH2PO4, and 10 d-glucose (pH 7.4, set by aeration with 95% O2 and 5% CO2). Coronal brain slices (thickness of 300 or 400 μm) containing the hippocampus were cut in ice-cold standard or modified ACSF with the use of a Leica VT1200S vibratome (Leica, Heidelberg, Germany). The slices were then transferred immediately to a nylon net submerged in standard ACSF for at least 40 min at a controlled temperature of 35°C (for patch-clamp experiments) or at room temperature (for extracellular recordings). After subsequent incubation for at least 1 h at room temperature, hemi-slices were transferred to the recording chamber, which was perfused with standard ACSF at a constant flow rate of ~2 ml/min. For all recordings, the temperature of the bath was maintained at 33°C.

**EXTRACELLULAR RECORDING OF iEPSPs**

Recordings of field excitatory postsynaptic potentials (fEPSPs) were obtained from the stratum radiatum of the CA1 region of the hippocampus after stimulation of the Schaffer collateral afferents. Extracellular recording electrodes were prepared from borosilicate capillaries with an internal filament and an outer diameter of 1.5 μm (Sutter Instruments, Novato, CA, USA) and were filled with 4 M NaCl (resistance, 1–2 MΩ). For stimulation of afferents, a concentric bipolar electrode (FHC, Bowdoin, ME, USA) was positioned ~300 μm from the recording site. Responses were triggered digitally every 20 s with the use of an interval generator (Master 8, FHC) and a stimulus isolator by application of a constant current pulse of 0.2–0.4 mA with a duration of 60 μs, which yielded a half-maximal response. For determination of the input–output (I–O) relation, the stimulation current was adjusted from 0 to 1 mA in steps of 0.1 mA. The fEPSPs were amplified with the use of an Axoclamp 2B amplifier (Axon Instruments, Union City, CA,
US), digitized, and then analyzed with Clampfit 9.02 software (Axon Instruments). Several kinetic parameters of fEPSPs were analyzed, but the slope values were considered for quantitation of the responses. For elicitation of LTP, after 10 min of stable baseline recording of fEPSPs evoked every 20 s at the current intensity that triggered 50% of the maximal fEPSP response, high-frequency stimulation (HFS) consisting of a single train of 100 stimuli at 250 Hz was delivered and recording was then continued for 60 min with stimulation of fEPSPs every 20 s.

WHOLE-CELL PATCH-CLAMP RECORDINGS

Whole-cell recordings from hippocampal CA1 pyramidal neurons were performed as previously described (Sanna et al., 2011). Recording pipettes were prepared from borosilicate glass with the use of a Flaming Brown micropipette puller (Molecular Devices, Union City, CA, USA). Resistance of the pipettes ranged from 2.5 to 4.5 MΩ when they were filled with an internal solution containing 135 mM potassium gluconate, 10 mM MgCl₂, 0.1 mM CaCl₂, 1 mM EGTA, 10 mM Heps–KOH (pH 7.3), and 2 mM ATP (disodium salt). We analyzed only recordings with access resistance of <25 MΩ (values usually ranged from 9 to 20 MΩ). Series resistance was not compensated, and cells were excluded from further analysis if access resistance changed by >20% during the course of the recording. Membrane potentials were recorded with the use of an Axopatch 200-B amplifier (Axon Instruments), filtered at 2 kHz, and digitized at 5 kHz. For patch-clamp experiments, we used pClamp 9.2 software (Molecular Devices, Union City, CA, USA), which allowed us to measure various characteristics of the neuronal membrane and action potentials (APs). For current-clamp experiments, we applied a protocol with injected current steps of 400-ms duration and ranging in intensity from −80 to 200 pA in intervals of 20 pA in order to hyperpolarize or depolarize the membrane potential. The parameters analyzed included resting membrane potential, input resistance, AP threshold, minimum injected current capable of evoking the first AP, spike latency (time required for the first AP to occur in response to depolarization), and spike frequency. Input resistance was calculated from only the hyperpolarizing current steps.

RECORDING OF SPONTANEOUS LOCOMOTOR ACTIVITY

Group-housed and SI mice were tested at the end of the 6-week housing period. To determine general locomotor activity levels and exploration habits, we used a motility meter (Omnitech Electronics Inc.). Animals were left in the same room in which the apparatus was placed for at least 2 h before the beginning of experiments in order to allow their habituation to the environment. Individual mice were placed in the center of a square arena (20.3 by 20.3 cm) and allowed to move freely for 60 min while being tracked by the automated tracking system. Parameters were monitored every 5 min (total of 12 acquisitions). Data obtained during the first 10 min were used for comparison among groups. The arena was assembled with specially designed sound-attenuating shells made of polypropylene and an expanded PVC sheet. The animals were isolated from noise of the recorder and printer used to acquire the data by placing these devices in a different room. The parameters measured included horizontal activity (number of photobeam interruptions), total distance traveled (centimeters), locomotion time (seconds), and rest time (seconds).

STATISTICAL ANALYSIS

Data are presented as means ± SEM and were compared by one-way analysis of variance (ANOVA) or Student’s t-test with the use of Prism software (version 5, GraphPad). A p value of <0.05 was considered statistically significant.

RESULTS

EFFECTS OF SI ON NEURONAL EXCITABILITY AND LTP INDUCTION IN THE HIPPOCAMPAL CA1 REGION OF C57BL/6J MICE

To examine the effect of SI on neuronal excitability in the CA1 region of the mouse hippocampus, we generated I–O curves by stimulating the Schaffer collateral glutamatergic afferents with increasing (0–1.0 mA) current intensity and recorded fEPSPs from the dendritic region within the CA1 stratum radiatum. Consistent with previous observations (Bartesaghi, 2004; Sanna et al., 2011), the intensity of the stimulatory current that evoked a half-maximal response (quantified by analysis of the fEPSP slope) was significantly (p < 0.05) lower in hippocampal slices from SI mice (0.45 ± 0.02 mA) compared with that for GH mice (0.37 ± 0.01 mA; Figure 1A).

We then examined the effect of SI on LTP induction in the CA1 region. After a 10-min baseline recording was obtained by stimulation at the current intensity that elicited a half-maximal response and at a frequency of 0.05 Hz, LTP was induced by HFS of the Schaffer collateral afferents with a single train of 100 stimuli of the same intensity and at 250 Hz. Separate experiments in hippocampal slices obtained from GH mice revealed that the LTP induced with such stimulus frequency was dependent on N-methyl-d-aspartate (NMDA) receptors, as bath perfusion of AP5 (50 µM) completely prevented its induction, and it was not influenced by the L-type voltage-dependent Ca²⁺ channel antagonist nifedipine (10 µM; Figure 1B). As expected (Sanna et al., 2011), the extent of LTP, calculated by averaging the slope values of fEPSPs recorded between 50 and 60 min after HFS, was significantly (p < 0.05) lower in slices from SI mice than in those from GH animals (Figures 1C,D). The effect of HFS on the I–O curves also differed between the two groups of animals. Indeed, at 60 min after HFS, the maximal response was enhanced to a greater extent in GH mice (+167%) than in SI animals (+60%; Figures 1E,F). The current intensity that evoked a half-maximal response after HFS did not differ significantly between GH and SI animals (data not shown).

ENHANCED SENSITIVITY OF SI ANIMALS TO THE INHIBITORY EFFECT OF ETOH ON LTP IN THE CA1 REGION

Given that SI was previously shown to increase the sensitivity of rats to the steroidogenic effect of acute systemic administration of EtOH (Serra et al., 2006), we examined the effect of EtOH on LTP in hippocampal slices from both GH and SI mice. The concentration of EtOH studied (40 mM) was selected as the highest level that had no effect on LTP in GH mice. EtOH (40 mM) was applied to hippocampal slices of both GH and SI mice 30 min before HFS and was found to significantly (p < 0.05) reduce (by 67%) the extent of LTP in SI mice but not in GH animals.
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FIGURE 1 | Effects of social isolation on neuronal excitability and LTP induction in the hippocampus of C57BL/6J mice. Field excitatory postsynaptic potentials (fEPSPs) were recorded in the dendritic CA1 region of hippocampal slices obtained from group-housed (GH) or socially isolated (SI) animals. (A) Input–output (I–O) relations determined by measurement of fEPSP slope in response to stimulation of Schaffer collaterals with current of increasing intensity (from 0 to 1.0 mA). Data are expressed as a percentage of the corresponding maximal response and are means ± SEM (n = 12–20). (B) LTP induction calculated between 50 and 60 min after HFS stimulation is not affected by the voltage-dependent calcium channel antagonist nifedipine (10 μM), but it is completely prevented by the perfusion (before and after HFS) of the NMDA receptor antagonist AP5 (50 μM). (C) LTP was induced in the CA1 region by HFS (arrow) of the Schaffer collateral–CA1 pathway. Data represent fEPSP slope at the indicated times expressed as a percentage of that at baseline and are means ± SEM (n = 8–12 slices). Above the bar graph, representative traces of fEPSPs obtained from GH and SI mice before (black trace) and 50 min after (red trace) high-frequency stimulation (HFS). (D) The extent of LTP was calculated by averaging the change in fEPSP slope apparent between 50 and 60 min after HFS. Data are means ± SEM (n = 8–12 slices). *p < 0.05 vs. GH mice. (E,F) I–O relations before and 60 min after HFS in GH (E) and SI (F) mice. Data are expressed as a percentage of the maximal response measured before HFS and are means ± SEM (n = 8–12 slices).

The ability of EtOH to inhibit LTP in the hippocampal CA1 region was previously shown to be dependent on stimulation of the local biosynthesis and release of 5α-reduced pregnane neurosteroids such as 3α,5α-THP (Izumi et al., 2007). To test whether such a mechanism was also operative under our experimental conditions, we examined the effect of finasteride, an inhibitor of 5α-reductase (Finn et al., 2006) that prevents the formation of GABAergic neurosteroids (Sanna et al., 2004). The co-application of EtOH (40 mM) and finasteride (1 μM) prevented the inhibitory effect of EtOH on LTP in hippocampal slices from SI mice, but it had no effect on LTP in those from GH animals (Figures 2A–C,E–G). In addition, finasteride application did not alter basal LTP when measured in slices from either GH or SI mice (results not shown).

Evaluation of the I–O relations before and after HFS, either in the absence or presence of EtOH or finasteride, revealed that the maximal post-HFS response was markedly reduced by EtOH in SI mice, but not in GH mice, and that finasteride abolished this...
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FIGURE 2 | Differential effects of EtOH and finasteride on LTP induction in the CA1 region of SI and GH mice. (A,E)

Representative traces of fEPSPs obtained from hippocampal slices of GH (A) and SI (E) mice before (black trace) and 60 min after (red trace) HFS in the absence or presence of EtOH or finasteride (Fin). EtOH (40 mM) was applied to hippocampal slices 30 min before HFS, whereas finasteride (1 μM) was applied 10 min before EtOH. (B,F) LTP was induced in the CA1 region of hippocampal slices from GH (B) and SI (F) mice by HFS (arrow) of the Schaffer collateral–CA1 pathway in the absence or presence of EtOH or finasteride, as indicated. Data represent fEPSP slope at the indicated times expressed as a percentage of that at baseline and are means ± SEM (n = 5 or 6 slices). (C,G) The extent of LTP was calculated by averaging the change in fEPSP slope apparent between 50 and 60 min after HFS in hippocampal slices from GH (C) and SI (G) mice. Data are means ± SEM (n = 5 or 6 slices).

*p < 0.05 vs. control. (D,H) I–O relations determined before and 60 min after HFS in GH (D) and SI (H) mice. Data are expressed as a percentage of the maximal response measured before HFS and are means ± SEM (n = 8–12 slices).

inhibitory effect of EtOH in slices from SI mice without having an effect in slices from GH animals (Figures 2D,H).

EFFECTS OF SI ON MEMBRANE EXCITABILITY OF CA1 PYRAMIDAL NEURONS

To evaluate further the effects of SI on CA1 neuronal excitability, we performed patch-clamp experiments in the current-clamp mode with single pyramidal neurons present in hippocampal slices of both GH and SI animals. The resting membrane potential was similar in SI (−65.7 ± 1.2 mV, n = 25 cells) and GH (−69.6 ± 1.3 mV, n = 37) mice (Figure 3A). We measured the input resistance by injecting hyperpolarizing current pulses at various intensities and recording the relative negative deflections in membrane potential. Analysis of the voltage–current curves indicated that input resistance did not differ significantly between GH and SI mice (GH, 0.221 ± 0.01 GΩ, n = 37; SI, 0.199 ± 0.02 GΩ, n = 25; Figures 3B,C). Firing of APs was also analyzed by injection of depolarizing current pulses (Figure 3D).
FIGURE 3 | Effects of SI on hippocampal CA1 pyramidal neuron excitability and their reversal by progesterone treatment. 

(A–C) Lack of effect of SI or progesterone (Prog) treatment on membrane resting potential (A) the membrane potential–current relation (B) or membrane input resistance (C) determined by whole-cell patch-clamp analysis in the current-clamp mode. SI mice were treated daily with progesterone or vehicle before analysis. Data are for 37, 25, and 9 neurons in (A) and for 37, 25, and 9 neurons in (B,C) for GH, SI, and progesterone-treated SI mice, respectively; those in (B,C) are means ± SEM.

(D) Representative membrane voltage responses to negative (−20 pA) and positive (+60 pA) current pulses applied to CA1 pyramidal neurons in hippocampal slices from GH or SI (with or without progesterone treatment) mice. 

(E–H) Effects of SI and progesterone treatment on the mean action potential (AP) threshold (E), minimum current intensity required for induction of the first AP (F), AP frequency at each depolarizing current step (G), and AP latency (H). Data are means ± SEM for 37 neurons of GH mice, 25 neurons of SI mice, and 9 neurons of SI mice treated with progesterone. *p < 0.01 vs. GH.

The threshold membrane potential for AP firing did not differ between the two groups of mice (Figure 3E). However, the minimum current intensity required for generation of an AP was significantly greater for CA1 pyramidal neurons of SI mice than for those of GH mice (Figure 3F). In addition, pyramidal neurons from SI mice were characterized by a reduced AP frequency (Figure 3G) and an increased AP latency (Figure 3H) compared with GH animals.

EFFECTS OF PROGESTERONE TREATMENT ON SI-INDUCED CHANGES IN NEURONAL EXCITABILITY AND LTP

Social isolation in rodents is associated with reduced plasma and brain levels of neuroactive steroids (Serra et al., 2000). We tested whether such an SI-induced decrease in neuroactive steroid levels might affect hippocampal neuronal excitability and synaptic plasticity. We thus treated a separate group of mice with progesterone (5 mg/kg, subcutaneous, once a day) during the entire 6-week period of SI (Figure 3). Progesterone treatment resulted in a partial reversal of the decrease in CA1 pyramidal neuron excitability apparent in SI mice, as reflected by the change in spike frequency (Figure 3G, n = 9), spike latency (Figure 3H, n = 9), and the minimum current required to evoke the first AP (Figure 3F, n = 9). Progesterone treatment also partially reversed the reduction in the extent of LTP induced by SI (Figure 4); this effect is particularly evident for the late phase of LTP, while the early phase appears to be less influenced by such treatment. Furthermore, given that we recently showed that SI results in a decrease in spontaneous locomotor activity in mice subjected to the open field test (Sanna...
We have found that rearing of mice in isolation for 6 weeks after weaning resulted in a decrease in the excitability of CA1 pyramidal neurons that was accompanied by a reduction in the extent of LTP induction as well as an enhanced inhibitory effect of EtOH on LTP as compared with control mice housed in groups. These effects of SI on neuronal excitability and LTP appear to be related to the decrease in neuroactive steroid levels induced by this condition, given that they were reversed by treatment of mice with progesterone during the SI period.

Postweaning SI in C57BL/6J mice is studied as an animal model of prolonged mild stress and has been shown to be associated with marked changes in the activity of the HPA axis, increased sensitivity to the steroidogenic effect of EtOH, and increased voluntary consumption of EtOH (Matsumoto et al., 1999; Serra et al., 2000, 2006; Sanna et al., 2011). We have now shown that SI-induced a marked decrease in neuronal excitability in CA1 pyramidal neurons of C57BL/6J mice. Current-clamp recordings thus revealed different responses of neurons from GH or SI mice to depolarizing current pulses injected to trigger the firing of APs. CA1 neurons of SI mice needed higher current pulses to evoke the first AP, and they showed a lower AP frequency and increased spike latency. Neither resting membrane potential nor input resistance differed significantly between CA1 neurons from the two types of mice. These data are consistent with those of previous studies showing that SI is associated with pronounced changes in neuronal membrane excitability (Moyer et al., 1996, 2000; Thompson et al., 1996; Tombaugh et al., 2005). Extracellular recordings of fEPSPs in the hippocampal CA1 region of SI mice also revealed a reduced excitability at the Schaffer collateral–CA1 glutamatergic synapses, as revealed by a rightward shift in the I–O curve compared with that of GH mice. These results are also in agreement with those of a previous study of the guinea pig hippocampus (Bartesaghi, 2004).

As we suggested previously (Sanna et al., 2011), the reduced excitability of CA1 pyramidal neurons of SI mice might be responsible for the reduced level of LTP induced at Schaffer collateral–CA1 synapses in these animals. Our present data are thus consistent with previous studies suggesting that long-term synaptic plasticity in several brain areas is markedly modified by the stress associated with SI (Roberts and Greene, 2003; Bianchi et al., 2006; Conrad et al., 2011).

We found that EtOH at 40 mM inhibited the induction of LTP in the CA1 region of SI mice but not in that of GH mice, consistent with the enhanced sensitivity to this drug previously shown to be induced by SI. The inhibitory effect of EtOH on LTP was previously shown to be dependent on its stimulation of neurosteroidogenesis (Izumi et al., 2007). Co-application of the 5α-reductase inhibitor finasteride was also previously shown to abolish the inhibitory effect of EtOH on GABAAR function as well as neurosteroid synthesis in the brain (Sanna et al., 2004). Ethanol impairs LTP or other forms of synaptic plasticity in several brain regions including the hippocampus (Morrisett and Swartzwelder, 1993; Izumi et al., 2005), striatum (Yin et al., 2007), and cerebellum (Belmeguenai et al., 2008), but a role for neurosteroids in the EtOH induced impairment of LTP in the rat hippocampus was only recently proposed (Izumi et al., 2007). The reduced levels of hippocampal excitability and LTP observed in SI mice might explain the deficits in learning and memory induced by isolation in C57BL/6J mice (Voikar et al., 2005) as well as in rats (Lu et al., 2003; Bianchi et al., 2006; Quan et al., 2010).
Table 1 | Changes in spontaneous locomotor activity in SI mice: effect of treatment with progesterone and co-treatment with progesterone and finasteride.

| Parameter                | Group-housed | Isolated | Isolated + progesterone | Isolated + progesterone + finasteride |
|--------------------------|--------------|----------|-------------------------|---------------------------------------|
| Horizontal activity      | 1607 ± 88.9  | 1266 ± 83.5* | 1479 ± 89.7            | 1190 ± 45.3*                          |
| Total distance (cm)      | 399.1 ± 38.76| 256.9 ± 272* | 332.1 ± 24.4           | 231 ± 20.5*                           |
| Locomotion time (s)      | 53.7 ± 7.52  | 34.3 ± 4.6*  | 37.8 ± 2.7*            | 35 ± 5.9*                             |
| Rest time (s)            | 242.2 ± 77   | 265.5 ± 7.3* | 260.8 ± 2.8*           | 261 ± 6.0*                            |

A group of SI mice were treated with progesterone (5 mg/kg, s.c.) or progesterone together with finasteride (25 mg/kg, s.c.) once a day for the 6-week isolation period. Data are means ± SEM for six mice and were obtained during the first 10 min (of a total of 60 min) of the experiment. *p < 0.05 vs. GH mice.

To examine whether the reduced brain levels of neurosteroids such as 3α,5α-THP and 3α,5α-THDOC associated with SI might be important for the changes in hippocampal excitability and in sensitivity to the steroidalogenic effect of EtOH induced by isolation in mice, we administered progesterone daily during the 6-week period of isolation in an attempt to restore the normal levels of neuroactive steroids in the hippocampus (Costa et al., 1995; Moran et al., 1998). We found that such progesterone treatment during SI partially reversed the decreases in the levels of neuronal membrane excitability and LTP induced by SI. It should be noted that the protective effect of progesterone treatment on SI-induced decreased in LTP appears to be mostly involving the late phase of this phenomenon, whereas the early phase was less influenced (Figure 4); however, the reason for such differential action is at present unknown.

Progesterone treatment also attenuated the decrease in spontaneous locomotor activity shown to be induced by SI in previous studies (Valzelli et al., 1974; Voikar et al., 2005; Fone and Forkess, 2008; Pietropaolo et al., 2008; Arndt et al., 2009; Sanna et al., 2011). Furthermore, the effects of progesterone on LTP and spontaneous locomotor activity induced by SI were prevented by the concomitant administration of finasteride.

Social isolation is associated with increased expression of several subunits of GABA<sub>A</sub>Rs, in particular α and γ subunits, in both rats (Serra et al., 2006) and mice (Sanna et al., 2011). Tonic inhibition is thought to play a key role in regulation of membrane excitability under both physiological and pathological conditions that have been shown to be associated with marked modulation of extrasynaptic GABA<sub>A</sub>R function and subunit expression (Maguire et al., 2005; Serra et al., 2006; Maguire and Mody, 2008; Sanna et al., 2009, 2011). We propose that the effects of SI on CA1 excitation and LTP are mediated in part by a decrease in excitability of granule cells in the dentate gyrus that results from an increase in GABAergic tonic current (Sanna et al., 2011) and which may lead to suppression of the activity of the entire dentate gyrus–CA3–CA1 circuitry. In line with this idea, Bartesaghi (2004) demonstrated that early isolation in guinea pigs results in a reduction in the synaptic function of the DG–CA3–CA1 neuronal circuitry. However, the impact of prolonged stress on hippocampal function may be more complex. In fact, Airan et al. (2007) working in the ventral hippocampal found that following chronic mild stress exposure in adult female rats there was diminished inflow through dentate, consistent with our results, but output from area CA1 was actually increased, resulting in an I–O mismatch.

Our present results provide new insight into changes in hippocampal function induced by SI in mice, including impairment of neuronal excitability and LTP associated with an increased sensitivity to the steroidalogenic effect of moderate doses of EtOH. Furthermore, the increased sensitivity to the neurosteroidogenic effect of EtOH associated with SI likely accounts for the greater inhibitory effect of EtOH on LTP in SI mice. These effects of SI may be related to the deficits in learning and memory and changes in responses to EtOH that are associated with isolation, and they warrant further investigation with regard to the control of GABAergic transmission by neurosteroids at the level of the dentate gyrus.

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