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なし

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2. 実用新案登録

なし

3.その他

なし

平成20～22年度　3年間　刊行物一覧

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Social isolation rearing-induced impairment of the hippocampal neurogenesis is associated with deficits in spatial memory and emotion-related behaviors in juvenile mice

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Abstract

Experiences during brain development may influence the pathogenesis of developmental disorders. Thus, social isolation (SI) rearing after weaning is a useful animal model for studying the pathological mechanisms of such psychiatric diseases. In this study, we examined the effect of SI on neurogenesis in the hippocampal dentate gyrus (DG) relating to memory and emotion-related behaviors. When newly divided cells were labeled with 5-bromo-2′-deoxyuridine (BrdU) before SI, the number of BrdU-positive cells and the rate of differentiation into neurons were significantly decreased after 4-week SI compared with those in group-housed mice. Repeated treatment of fluoxetine prevented the SI-induced impairment of survival of newly divided cells and ameliorated spatial memory impairment and part of aggression in SI mice. Furthermore, we investigated the changes in gene expression in the DG of SI mice by using DNA microarray and real-time PCR. We finally found that SI reduced the expression of development-related genes *Nurr1* and *Npas4*. These findings suggest that communication in juvenile is important in the survival and differentiation of newly divided cells, which may be associated with memory and aggression, and raise the possibility that the reduced expression of *Nurr1* and/or *Npas4* may contribute to the impairment of neurogenesis and memory and aggression induced by SI.

Keywords: aggressive behavior, fluoxetine, learning and memory, neurogenesis, social isolation.
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Experiences during a critical period of brain development may affect structural and functional development and maturation of the brain (Mataga *et al.* 2004; Mirescu *et al.* 2004), and influence behavior including cognitive and emotional functions which could be attributable to the expression or exacerbation of developmental disorders (Castellanos and Tannock 2002). Rearing animals in isolation is a relevant paradigm to investigate the effect of early life stress and for understanding the pathogenesis of certain neurological and psychiatric diseases (Myhrer 1998; Whitaker-Azmitia *et al.* 2000). Behavioral changes induced by isolation rearing have been characterized, including

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Abbreviations used: 5-HT, 5-hydroxytryptamine; BrdU, 5-bromo-2′-deoxyuridine; DG, dentate gyrus; Flx, fluoxetine; GCL, granule cell layer; GFAP, glial fibrillary acidic protein; GH, group-housed; NeuN, neuronal nuclei; Npas4, neuronal PAS domain 4;Nurr1, nuclear receptor subfamily 4, group A, member 2; PBS, phosphate-buffered saline; RT, reverse transcription; Sal, saline; SGZ, subgranular zone; SI, social isolation; SSRI, selective serotonin reuptake inhibitors.

enhanced locomotor activity under a novel environment (Wilkinson *et al.* 1994), aggressive behaviors (Wongwitdecha and Marsden 1996), and impairment of pre-pulse inhibition (Day-Wilson *et al.* 2006) and spatial learning in a water maze test (Lu *et al.* 2003).

The social environment in early life significantly influences not only the organization of behavior but also neurochemical development of the brain. For instance, dopamine and serotonin systems are affected by social isolation (SI) in the nucleus accumbens (Hall *et al.* 1998), prefrontal cortex (Heidbreder *et al.* 2000) and hippocampus (Muchimapura *et al.* 2003). The neuroanatomic consequences of isolation rearing include decreased spine density of pyramidal neurons in the prefrontal cortex and hippocampus (Silva-Gomez *et al.* 2003) and fewer hippocampal synapses (Varty *et al.* 1999).

Hippocampal development is affected by environmental factors, but the underlying mechanisms are unclear. Accumulating evidence has demonstrated that neurogenesis occurs in adults in certain brain areas such as the hippocampus, in which newly divided neurons play a role in physiological function (Lledo *et al.* 2006). Recent studies suggested that the impairment of adult neurogenesis is involved in the development and expression of neuropsychiatric disorders (Jacobs *et al.* 2000; Reif *et al.* 2006; Maeda *et al.* 2007). For instance, the genesis of stem-like cells in the dentate gyrus (DG) of the hippocampus is decreased in patients with schizophrenia, which may contribute to the pathogenesis of the disorder (Reif *et al.* 2006).

It has been demonstrated that some mood-stabilizing drugs and selective serotonin reuptake inhibitors (SSRI) enhance adult neurogenesis in the hippocampus, and the effect may contribute to their clinical effects (Santarelli *et al.* 2003; Encinas *et al.* 2006). For example, an SSRI fluoxetine (Flx) prescribed for depression and anxiety disorders including obsessive compulsive disorder and panic disorder is reported to enhance neurogenesis in the hippocampus (Santarelli *et al.* 2003; Encinas *et al.* 2006).

It is well known that some genes, such as *reelin* and *brain-derived neurotrophic factor* regulate the development and migration of newly divided cells (Polleux *et al.* 2002; Gong *et al.* 2007). They are supposed to be associated with cognitive deficits in mental disorders (Angelucci *et al.* 2005; Fatemi 2005). It remains to be determined whether gene expression in the hippocampus is affected by environmental stress (e.g., SI) in early life.

In the present study, to investigate the effects of early life stress on neurogenesis in the hippocampus and on cognitive function and emotion related-behaviors, mice after weaning were subjected to SI rearing for 4 weeks. In addition, we investigated the effect of Flx on SI-induced impairment of survival of newly divided cells in the hippocampus, and emotion related-behaviors and memory impairment. Finally, to identify the genes whose expression in the DG of hippocampus is affected by SI, we examined the changes in gene expression in SI mice by using DNA microarray and real-time reverse transcription (RT)-PCR.

Materials and methods

Animals

The Institute for Cancer Research mice (Japan SLC Inc., Hamamatsu, Japan) were purchased when they were 3 and 8 weeks old and used for the experiments. The study was completed exclusively with male mice, because estrogen in female mice affects memory and SI-induced emotion-related behaviors (Li *et al.* 2004; Starkey *et al.* 2007). They were housed under a standard 12-h light/dark cycle (light phase 8:45 AM–20:45 PM) at a constant temperature of 23 ± 1°C with free access to food and water throughout the experiments. The animals were handled in accordance with the guidelines established by the Institutional Animal Care and Use Committee of Kanazawa University, the Guiding Principles for the Care and Use of Laboratory Animals approved by the Japanese Pharmacological Society and the National Institutes of Health Guide for the Care and Use of Laboratory Animals.

Isolation rearing

After 3 days of acclimatization, mice were randomly divided into two groups: SI rearing and group-housed (GH) rearing. Mice in the SI group were individually housed in wire-topped opaque polypropylene cages (20 cm× 12 cm× 10 cm) while other mice in the GH group continued to be housed under normal conditions (five per cage) in wire-topped clear plastic cages (34 cm× 22 cm× 15 cm). After the 4-week SI, mice were subjected to behavioral and histological analyses as described below. During the behavioral analysis, the housing conditions were maintained.

Drug administration

Both 5-Bromo-2′-deoxyuridine (BrdU) and Flx were purchased from Sigma-Aldrich (St Louis, MO, USA) and dissolved in saline. To label newly divided cells in the DG, BrdU (75 mg/kg) was injected intraperitoneally (i.p.) three times at 2 h intervals. Flx (10 mg/kg) or saline was administrated i.p. once a day for at least 2 weeks. Daily administration was started 2 weeks after SI and continued until the end of the study. During behavioral analysis, Flx was administrated 1 h and 30 min before water maze test and intruder-evoked aggressive test, respectively.

Immunohistochemistry

In histological analysis, mice were used without behavioral analyses. They were deeply anesthetized with diethyl ether at the indicated time and perfused transcardially with saline, followed by 4% paraformaldehyde in 0.1 M phosphate-buffered saline (PBS, pH 7.4). Their brains were removed, post-fixed in the same fixative and then cryoprotected. Thick coronal brain sections of 30 μm were cut on a cryostat and mounted on slides. Every fifth section was collected between stereotaxic coordinates bregma −1.2 to 3.0 according to the brain atlas (Paxinos and Franklin 2004). Sections were treated overnight with 0.1% nonidet-40/0.01 M PBS (pH 7.2) at 4°C and

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denatured in the microwave oven in 0.01 M citrate buffer (pH 6.0). After blocking in 10% goat serum/PBS with 0.1% nonident-40 for 30 min, BrdU-positive cells in the sections were detected using a BrdU labeling and detection kit 2 (Roche Diagnotics GmbH, Mannheim, Germany) according to the manufacturer's instructions.

For double-staining of BrdU/neuronal nuclei (NeuN, neuronal marker) and BrdU/glial fibrillary acidic protein (GFAP, astroglial marker), sections were pre-treated with 1 M HCl for 30 min at 37°C, followed by 10 min in 0.1 M borate buffer and then washed in PBS before blocking. Rat anti-BrdU antibody (1 : 200; Abcam, Cambridge, UK), mouse anti-NeuN antibody (1 : 100; Chemicon, Temecula, CA, USA) and mouse-GFAP antibody (1 : 1500; Sigma-Aldrich) diluted in PBS containing 0.1% Triton X-100 and 5% goat serum was applied to sections which were then incubated overnight at 4°C and for 6 h at around 25°C. After washing in PBS, goat anti-rat Alexa 568 and anti-mouse Alexa 488 (1 : 1000; Invitrogen, Eugene, OR, USA) were added to sections for 2 h at room temperature.

Quantification of immunostaining cells

Every fifth section throughout the hippocampus (total 12 sections from each mouse) was processed for BrdU immunohistochemistry. All BrdU-labeled cells in the subgranular zone (SGZ), hilus and granule cell layer (GCL) were assessed using a light microscope (Axio Imager; Zeiss, Jene, Germany) and counted by an experimenter blinded to the code. To distinguish single cells within clusters, all counts were performed at 400 × magnification (objective; 40 ×). To obtain the total number of cells per DG, we multiplied the counted number of positive cells by five (Maeda *et al.* 2007).

Double-stained cells were quantified using a confocal laser scanning microscope (LSM 510; Zeiss). Each cell was analyzed along the entire 'z' axis. Approximately 20 BrdU-positive cells in each mouse were randomly identified between five and six sections. Ratios of BrdU-positive cells co-labeled with NeuN or GFAP among BrdU-positive cells were determined.

Water maze test

After the 4-week SI, a water maze test was carried out as described previously (Jhoo *et al.* 2004; Miyamoto *et al.* 2005). Briefly, a pool (120 cm in diameter) was prepared, and the water temperature was maintained at 21–23°C. Swimming paths were analyzed by a computer system with a video camera (TAMRON, Saitama, Japan).

In the training trials, the platform (7 cm in diameter) was submerged 1 cm below the water surface. After reaching the platform, the mouse was allowed to remain on it for 20 s. If the mouse did not find the platform within 60 s, the trial was terminated and the animal was put on the platform for 20 s. After training trials for 6 days, mice were subjected to the probe test on day 7 where in they swam for 60 s in the pool without the platform. We measured the time spent in each quadrant of the pool as a measure of spatial memory. One hour after the probe test, to measure swimming ability or motivation, mice were subjected to the visible test in which the platform was marked with a flag that protruded 12 cm above the water surface to be highly visible, but in a new location. Three starting positions were used pseudo-randomly and each mouse was subjected to three trials per day in the training trials (day 1–6) and visible test (day 7). During training trials and the visible test, we measured both path length (swim distance) and escape latency as measures of performance.

Intruder-evoked aggressive test

We used 8-week-old male Institute for Cancer Research mice as intruders which have not shown aggressive behaviors against their peers. The day after the probe test in the water maze test (e.g., day 8), an intruder-evoked aggressive test was carried out as previously reported (Miczek and O'Donnell 1978). A resident mouse was habituated in a test cage (20 cm× 12 cm× 10 cm) for 10 min, and then an intruder mouse was put in the test cage. The investigating behavior performed by the resident mouse against the intruder was observed for 10 min. The frequency of attacking/biting and duration of aggression including attacking/biting, tail rattling, aggressive grooming, sideways posturing and pushing under were analyzed. The behavioral observation was made by the blinded experimenters.

Total RNA isolation for DNA microarray and real-time RT-PCR

Mice reared under GH and SI conditions for 3 days, 2 weeks, and 4 weeks were decapitated and their brains were removed. These mice had never been used for behavioral experiments. Brain slices including the hippocampus were made using brain matrix and the DG of the hippocampus was isolated using a dissecting microscope (AS ONE Co., Ltd., Osaka, Japan). Tissues from two animals and four animals were pooled as one sample for DNA microarray and real-time RT-PCR, respectively. Total RNA was isolated using the RNeasy Mini Kit (Qiagen, Hilden, Germany).

DNA microarray and expression profiling

The purified total RNA was checked for quality using Bioanalyzer 2100 electropherograms (Agilent; Santa Clara, CA, USA) and used for expression profiling with GeneChip mouse genome 430 2.0 arrays (Affymetrix; Santa Clara, CA, USA) containing 45 101 probe sets, according to the protocol provided by the manufacturer. Briefly, 5 μg of total RNA was reverse-transcribed into double-stranded cDNA with a T7-Oligo (dT) primer. Labeled cRNA was synthesized in the presence of T7 RNA polymerase and biotin-labeled nucleotides fragmented by metal-induced hydrolysis and hybridized overnight to the array. Each array was washed, stained with streptavidin-coupled phycoerythrin and scanned by a GCS3000 laser scanner (Affymetrix).

The resulting expression profiles were pre-processed using the robust multi-array average of G+C content (GCRMA) algorithm from Bioconductor (http://www.bioconductor.org/) in the statistical programming language R (http://www.r-project.org/). The changes in gene expression between GH and SI groups were quantified using a general linear model in the limma package from Bioconductor. Scatter plots and hierarchical clustering analysis were carried out in SPOTFIRE 8.2.1 (TIBCO Software Inc., Palo Alto, CA, USA).

Quantitative analyses of *Nurr1* and *Npas4* mRNA by real-time RT-PCR

Total RNA isolated from the DG was converted into cDNA using SuperScript III First-Strand Synthesis System for RT-PCR (Invitrogen). Levels of mRNA expression were quantified by using a 7300 real-time PCR System (Applied Biosystems, Foster City, CA, USA). The quantitative real-time PCR was performed in a volume of 25 μL with 500 ng of cDNA and 500 nM primers in the Power SYBR Green Master Mix (Applied Biosystems). The primers used were as follows: 5′-ATGACCAGCCTGGACTATTCC-3′ (forward) and 5′-CAGGAGATCGTAGAACTGCTGGA-3′ (reverse) for *Nurr*1 and

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5′-AGCATTCCAGGCTCATCTGAA-3′ (forward) and 5′-GGCGA AGTAAGTCTTGGTAGGATT-3′ (reverse) for *Npas*4. The mouse glyceraldehyde-3-phosphate dehydrogenase was used as an internal control (Applied Biosystems).

Statistical analysis

All data were expressed as the mean ± SE. Differences between two groups were analyzed by two-tailed Student's *t*-test or chi-square (χ^2) test. Differences among multigroups were analyzed by ANOVA followed by the Bonferroni's test when *F* ratios were significant ($p < 0.05$). Differences among multigroups of path length in the water maze test were analyzed by ANOVA with repeated measures, followed by the Bonferroni's test when *F* ratios were significant ($p < 0.05$).

Results

Effect of SI after weaning on newly divided cell proliferation in the DG of the hippocampus

To examine the effect of SI after weaning on newly divided cell proliferation in the hippocampus, BrdU was injected on the last day of the 4-week isolation and the number of BrdU-labeled cells was counted 24 h after the injection (Fig. 1a). As shown in Fig. 1b, BrdU-positive cells in SI and GH mice were observed as clusters, and there were no apparent differences in the morphology and location. Most of the BrdU-labeled cells were found in the SGZ of the DG in both GH and SI mice (Fig. 1b). There was no significant difference in the number of BrdU-labeled cells in the hippocampus between SI and GH mice (Fig. 1c).

Effect of SI after weaning on the cell survival of newly divided cells in the DG of hippocampus

Next, to investigate the effect of SI after weaning on the survival of newly divided cells, BrdU was injected one day before starting the SI and the number of BrdU-labeled cells in the hippocampus was counted after the 4-week SI (Fig. 2a). As shown in Fig. 2b, there was an apparent difference between SI and GH mice in the number and location of BrdU-labeled cells in the DG of hippocampus. Some of the BrdU-labeled cells in GH mice were found in the GCL of DG whereas in SI mice, fewer cells were detected in the GCL. The total number of BrdU-labeled cells in the hilus + SGZ + GCL of SI mice was 75% of that in GH mice, and the number of BrdU-labeled cells in the GCL of SI mice was 63% of that in GH mice. The number of BrdU-labeled cells in the hilus + SGZ + GCL and GCL of hippocampus in SI mice was significantly decreased compared with that in GH mice (Fig. 2c).

Effect of SI after weaning on the differentiation of newly divided cells in the DG of hippocampus

To examine the effect of SI after weaning on the differentiation of newly divided cells, we counted NeuN- and GFAP-positive cells among BrdU-labeled cells in the hippocampus. Mice were subjected to SI after BrdU-labeling and killed after the 4-week SI for double-immunostaining (Figs 3a and 4a). Most of the BrdU-labeled cells in GH mice were NeuN-positive, but some were NeuN-negative (Fig. 3b). The rate of NeuN-positive cells among BrdU-labeled cells in the hilus + SGZ + GCL of SI mice was significantly lower than that in GH mice (Fig. 3c). The impairment of neural differentiation of BrdU-labeled cells was also evident in the SGZ of SI mice compared with GH mice, while there was no difference in the hilus and GCL. Thus, the rate of NeuN-positive cells among BrdU-labeled newly divided cells in the hilus + SGZ + GCL and SGZ of hippocampus in SI mice was significantly lower than that in GH mice (Fig. 3c). By contrast, a small fraction of

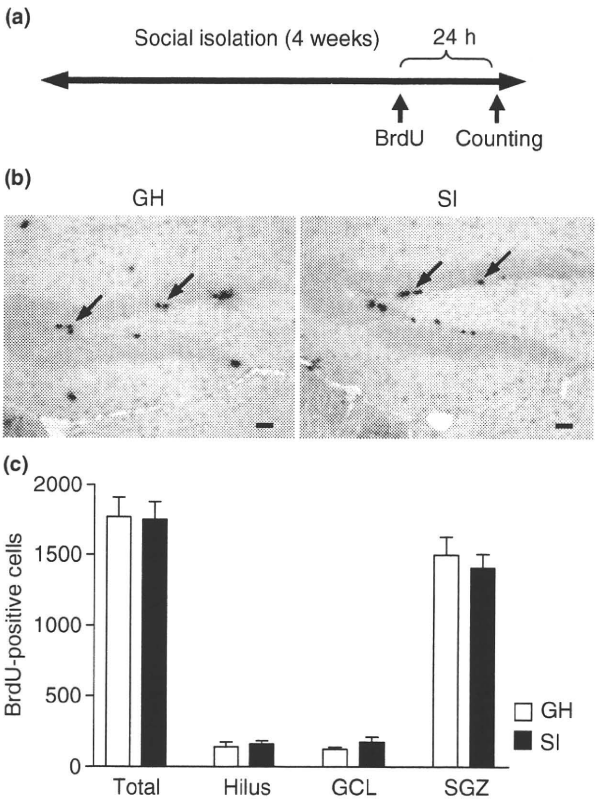

Fig. 1 Effect of social isolation (SI) for 4 weeks from 3-week-old in mice on the proliferation of newly divided cells in the dentate gyrus of hippocampus. 5-bromo-2′-deoxyuridine (BrdU; 75 mg/kg, i.p.) was injected three times at 2 h intervals on the last day of 4-week isolation. Animals were killed 24 h after the last injection of BrdU, and BrdU-positive cells in the subgranular zone (SGZ), hilus, and granule cell layer (GCL) were counted as described in *Materials and Methods*. (a) Experimental schedule. (b) Representative photographs showing the distribution of BrdU-positive cells in group-housed (left) and SI (right) mice, respectively. Scale bar: 200 μm. (c) Total numbers of BrdU-positive cells are expressed as the sum of the number in the SGZ (arrows), hilus, and GCL. Values indicate the mean ± SE ($n = 4$).

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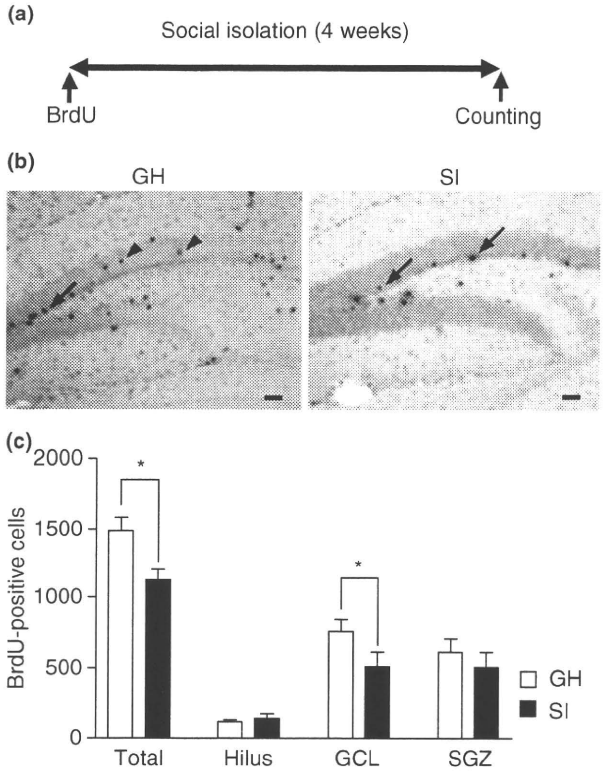

Fig. 2 Effect of social isolation (SI) for 4 weeks in 3-week-old mice on the survival of newly divided cells in the dentate gyrus of hippocampus. 5-bromo-2′-deoxyuridine (BrdU; 75 mg/kg, i.p.) was injected three times at 2 h intervals one day before starting 4-week isolation. Animals were killed after SI, and BrdU-positive cells in the subgranular zone (SGZ), hilus, and granule cell layer (GCL) were counted as described in *Materials and methods*. (a) Experimental scedule. (b) Representative photographs showing the distribution of BrdU-positive cells in group-housed (GH, left) and SI (right) mice, respectively. Scale bar: 200 μm. (c) Total numbers of BrdU-positive cells were expressed as the sum of the number in the SGZ (arrows), hilus, and GCL (arrowheads). Values indicate the mean ± SE (n = 7). *p < 0.05 versus GH (two-tailed t-test).

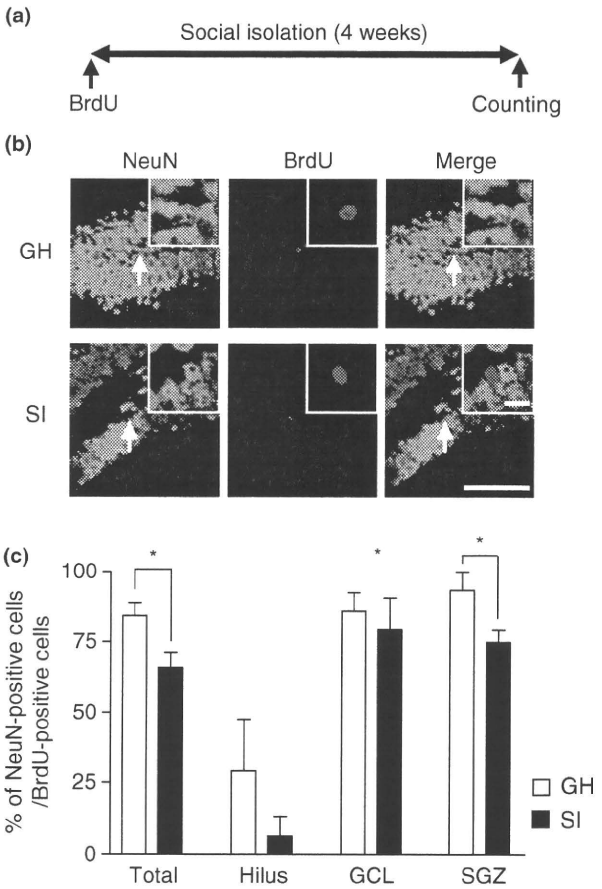

Fig. 3 Effect of social isolation (SI) for 4 weeks in 3-week-old mice on neurogenesis in the dentate gyrus. 5-bromo-2′-deoxyuridine (BrdU; 75 mg/kg, i.p.) was injected three times at 2 h intervals before 4-week isolation. Animals were killed after SI, and BrdU-labeled cells in the subgranular zone, hilus, and granule cell layer were counted as described in *Materials and Methods*. (a) Experimental scedule. (b) Representative photographs showing confocal analysis to determine the percentage of neurons [neuronal nuclei (NeuN)-positive cells: green] among the population of newly divided cells (BrdU-positive cells: red) at 4 weeks after BrdU labeling (double-stained cells: yellow). Scale bar: 100 μm (inset, 10 μm). (c) Percentage of neurons (NeuN-positive cells) among BrdU-positive cells. Values indicate the mean ± SE (n = 5). *p < 0.05 versus group-housed (two-tailed t-test).

BrdU-labeled cells was co-labeled with an astrocyte marker, GFAP (Fig. 4b). There was no significant difference between GH and SI mice in the rate of GFAP-positive cells among BrdU-labeled newly divided cells in the hilus + SGZ + GCL, hilus, SGZ or GCL of hippocampus (Fig. 4c).

Effect of repeated Flx treatment on SI-induced spatial learning and memory deficits

To examine the functional significance of SI-induced impairment of neurogenesis in the hippocampus, we compared the performance of GH and SI mice in the water maze test which was used to examine spatial learning and memory associated with the hippocampal function (Jhoo *et al.* 2004; Miyamoto *et al.* 2005). At the same time, we investigated the effect of repeated administration of Flx, an SSRI reported to enhance neurogenesis in the hippocampus (Malberg *et al.* 2000; Santarelli *et al.* 2003) on maze performance in GH and SI mice. Repeated daily administration of Flx at a dose of 10 mg/kg (Encinas *et al.* 2006) was started 2 weeks after starting the SI until the end of the maze test (Fig. 5a), as we confirmed that SI for 2 weeks after weaning had little effect on cell proliferation and survival of BrdU-labeled cells in the hippocampus [data not shown, p = 0.995 (hilus + SGZ + GCL)].

There was no difference in the time spent in each quadrant of the pool among four groups of mice in the pre-probe test that was carried out before training trials [data not shown, $F_{(3,38)} = 0.246$; $p = 0.864$], indicating that

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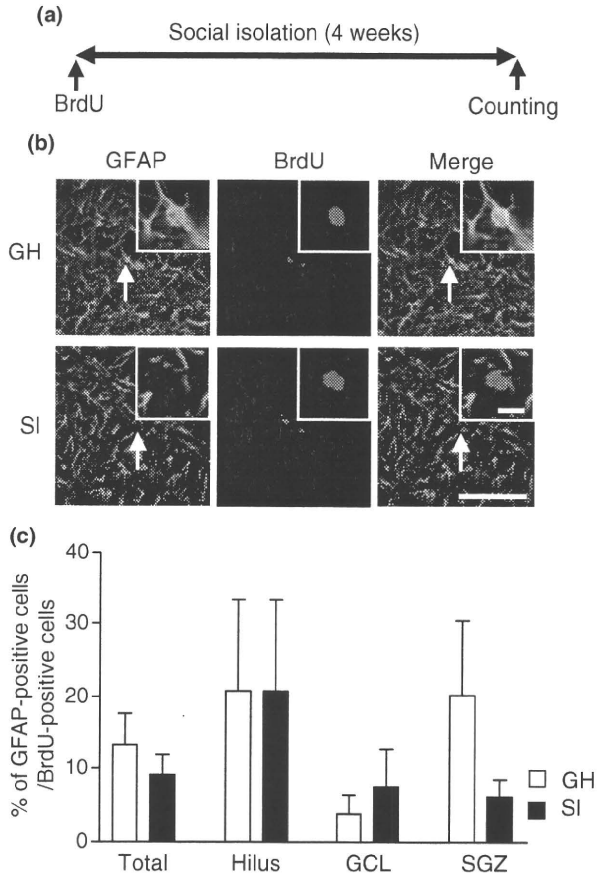

Fig. 4 Effect of social isolation (SI) for 4 weeks in 3-week-old mice on gliogenesis in the dentate gyrus. 5-bromo-2′-deoxyuridine (BrdU; 75 mg/kg, i.p.) was injected three times at 2 h intervals before 4-week isolation. Animals were killed after SI, and BrdU-labeled cells in the subgranular zone, hilus, and granule cell layer were counted as described in *Materials and Methods*. (a) Experimental design. (b) Representative photographs showing confocal analysis to determine the percentage of astroglial cells [glial fibrillary acidic protein (GFAP)-positive cells: green] among the population of newly divided cells (BrdU-positive cells: red) at 4 weeks after BrdU labeling (double-stained cells: yellow). Scale bar: 100 μm (inset, 10 μm). (c) Percentage of glial cells (GFAP-positive cells) among BrdU-positive cells. Values indicate the mean ± SE ($n = 4$).

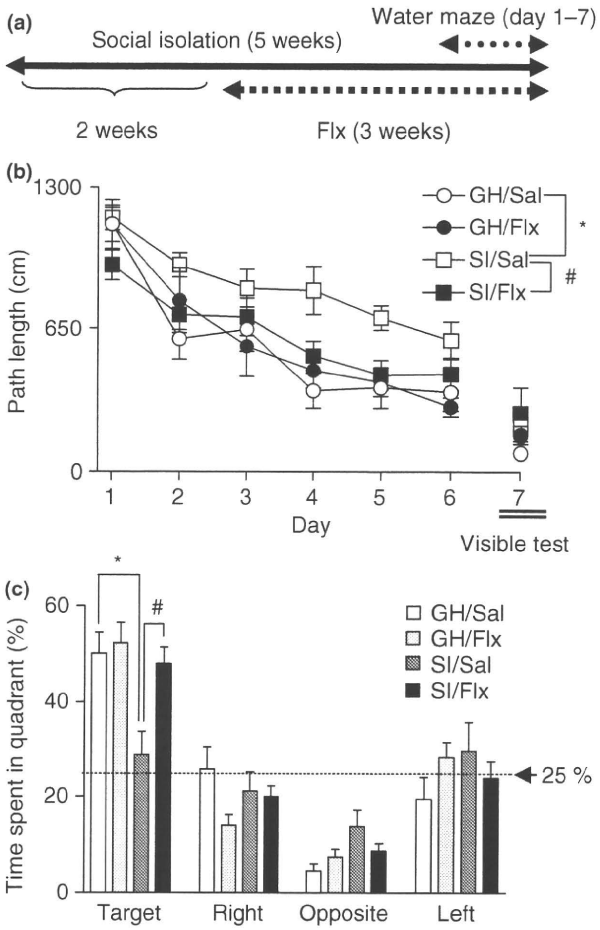

Fig. 5 Effect of repeated fluoxetine (Flx) treatment on social isolation (SI)-induced spatial learning and memory deficits in the water maze test. Mice were subjected to SI for 4 weeks and then to training trials on days 1–6, and visible and probe tests on day 7. The daily administration of Flx (10 mg/kg, i.p.) was started 2 weeks after SI and continued to the end of the water maze test. GH/Sal: saline-treated group-housed mice ($n = 7$); GH/Flx: fluoxetine-treated GH mice ($n = 10$), SI/Sal: saline-treated SI mice ($n = 12$), SI/Flx: fluoxetine-treated SI mice ($n = 10$). (a) Experimental design. (b) Path length (swim distance) to find the hidden platform during training trials (day 1–6) and visible test (day 7). (c) Time spent in quadrants in the probe test (day 7). Values indicate the mean ± SE. $^{*}p < 0.05$ versus GH/Sal, $^{\#}p < 0.05$ versus SI/Flx.

neither the housing condition nor Flx treatment had any effect on space preference before maze training.

We preliminarily checked that SI did not affect swim speed [data not shown, $F_{(1,8)} = 0.730$; $p = 0.418$]. Furthermore, previous paper suggested that treatment of repeated Flx (10 mg/kg) did not affect swim speed (Song *et al.* 2006). In addition to results of visible test in this study, these findings indicate that the changes in performance during the training and probe trials were not because of an impairment of swimming ability or motivation.

On the other hand, there was a significant difference in performance (path length) of four different groups of mice [saline-treated GH (GH/Sal) mice, Flx-treated GH (GH/Flx) mice, saline-treated SI (SI/Sal) mice, and Flx-treated SI (SI/Flx) mice] during training trials on days 1 to 6 (Fig. 5b, ANOVA with repeated measures: group, $F_{(3,38)} = 6.134$, $p < 0.05$; day, $F_{(5,185)} = 39.796$, $p < 0.001$; group × day, $F_{(15,185)} = 1.128$, $p = 0.334$). *Post-hoc* analysis indicated that performance by SI/Sal mice was significantly impaired compared with GH/Sal mice, suggesting that SI in juveniles induces the impairment of spatial learning. Furthermore, repeated Flx treatment significantly improved performance in SI mice although it had no effect on

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performance in GH mice. When escape latency was analyzed as a measure of performance, the same results were obtained [data not shown, $F_{(3,38)} = 7.217$; $p < 0.001$].

Next day (day 7), the animals were subjected to the probe test and then to the visible test. In the probe test (Fig. 5c), there was a significant difference in time spent in the target quadrant in which the submerged platform had been located during training trials among four groups [$F_{(3,38)} = 6.763$, $p < 0.05$]. *Post-hoc* analysis by Bonferroni's test indicated that SI/Sal mice spent significantly less time than GH/Sal mice, indicating an impairment of spatial memory. Furthermore, repeated Flx treatment significantly improved the impairment of spatial memory in SI mice, although the treatment had no effect on performance in GH mice. In the visible test conducted after the probe test on day 7, there was no significant difference in performance (both path length and escape latency) among the four groups of animals (Fig. 5b).

Effect of Flx on SI-induced aggressive behaviors

The day after the water maze probe test (day 8), the animals were used in the intruder-evoked aggressive test (Fig. 6a). Aggressive biting behavior was observed in all four groups (1/13 in GH/Sal, 0/9 in GH/Flx, 8/12 in SI/Sal, and 2/10 in SI/Flx mice). The difference between GH/Sal and SI/Sal groups was statistically significant ($\chi^2 = 9.420$, $p < 0.01$). Repeated Flx treatment in SI mice significantly reduced the rate of mice exhibiting biting ($\chi^2 = 4.791$, $p < 0.05$). Similarly, the rate of SI/Sal mice showing tail rattling against the intruder (8/12) was significantly increased compared to GH/Sal (2/13, $\chi^2 = 6.838$, $p < 0.01$). Repeated Flx treatment significantly reduced the rate of animals showing tail rattling in the SI group (1/10, $\chi^2 = 7.246$, $p < 0.01$) without affecting behavior in the GH group (0/9).

As shown in Fig. 6(b) and (c), there were significant differences in biting counts [Fig. 6b; $F_{(3,40)} = 3.650$, $p < 0.05$] and total time of aggression [Fig. 6c; $F_{(3,40)} = 16.075$, $p < 0.001$] among the four different groups of mice. *Post-hoc* analysis indicated that biting counts and total time of aggression against the intruder in SI/Sal were increased compared with those in GH/Sal mice. There was no difference in biting counts between SI/Sal and SI/Flx mice (Fig. 6b), but the total time of aggression in SI/Flx mice was significantly reduced compared to SI/Sal mice (Fig. 6c).

Effect of Flx on SI-induced reduction of survival of newly divided cell in the hippocampus

To examine the effect of repeated administration of Flx on SI-induced impairment of cell survival in the DG of the hippocampus, we compared the number of BrdU-positive cells in the DG among the four groups. Repeated daily administration of Flx at a dose of 10 mg/kg was started 2 weeks after starting SI and continued for 2 weeks (Fig. 7a).

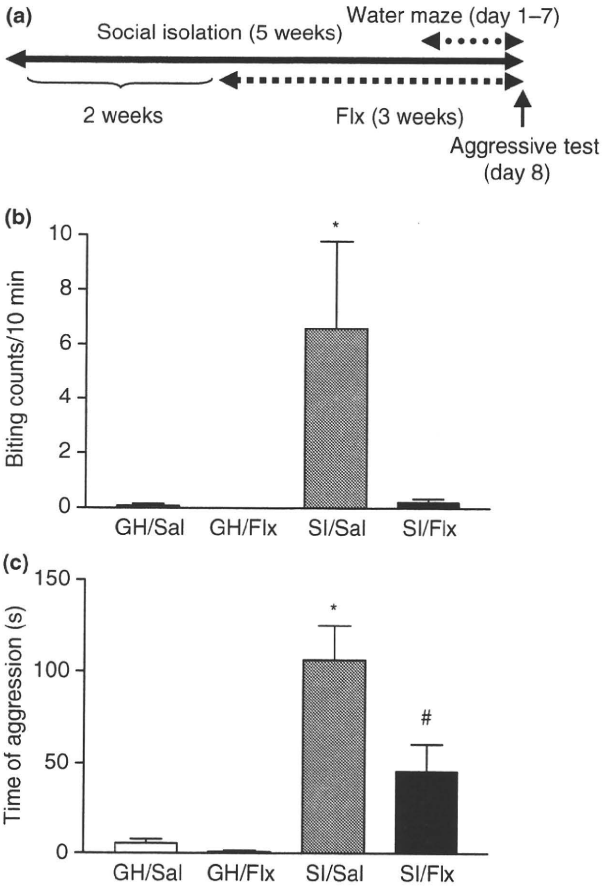

Fig. 6 Effect of repeated fluoxetine (Flx) treatment on social isolation (SI)-induced aggressive behavior in the intruder-evoked aggressive test. Mice that had previously been subjected to the water maze test following SI for 4 weeks were used in the intruder-evoked aggressive test. Daily administration of Flx (10 mg/kg, i.p.) was started 2 weeks after SI and continued to the end of the behavioral test. Frequency of biting and duration of aggressive behavior against the intruder were measured for 10 min. GH/Sal: saline-treated group-housed mice ($n = 13$), GH/Flx: fluoxetine-treated GH mice ($n = 9$), SI/Sal: saline-treated SI mice ($n = 12$), SI/Flx: fluoxetine-treated SI mice ($n = 10$). (a) Experimental schedule. (b) Biting counts. (c) Total time of aggression. Values indicate the mean ± SE. $^*p < 0.05$ versus GH/Sal, $^\#p < 0.05$ versus SI/Sal.

As shown in Fig. 7b, there was an apparent difference in the number and location of BrdU-labeled cells in the DG of the hippocampus in SI/Sal mice compared with GH/Sal, GH/Flx, and SI/Flx groups. ANOVA of the number of BrdU-positive cells in the GCL ($F = 6.183$, $p < 0.01$) and in the DG of hippocampus ($F = 6.536$, $p < 0.01$) revealed a significant effect of treatment. *Post-hoc* analysis with Bonferroni's test indicated that the total number of BrdU-positive cells in the DG of the hippocampus was significantly reduced in SI/Sal mice compared with GH/Sal mice ($p < 0.05$), and this impairment of survival of newly divided cells in the SI/Sal group was significantly ameliorated by

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