Social isolation–related depression accelerates ethanol intake via microglia-derived neuroinflammation

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Social isolation is common in modern society and is a contributor to depressive disorders. People with depression are highly vulnerable to alcohol use, and abusive alcohol consumption is a well-known obstacle to treating depressive disorders. Using a mouse model involving isolation stress (IS) and/or ethanol intake, we investigated the mutual influence between IS-derived depressive and ethanol-seeking behaviors along with the underlying mechanisms. IS increased ethanol craving, which robustly exacerbated depressive-like behaviors. Ethanol intake activated the mesolimbic dopaminergic system, as evidenced by dopamine/tyrosine hydroxylase double-positive signals in the ventral tegmental area and c-Fos activity in the nucleus accumbens. IS-induced ethanol intake also reduced serotonergic activity, via microglial hyperactivation in raphe nuclei, that was notably attenuated by a microglial inhibitor (minocycline). Our study demonstrated that microglial activation is a key mediator in the vicious cycle between depression and alcohol consumption. We also propose that dopaminergic reward might be involved in this pathogenicity.

INTRODUCTION
Depression is the most common neuropsychiatric disorder, with a prevalence of 350 million individuals worldwide (1). Patients with depressive disorder had a 20–40-fold higher risk of suicidality than healthy individuals (2). Recently, our society has emphasized individualism, and these social changes have led to an increase in the prevalence of depressive disorder (3, 4). Social withdrawal is a more potent cause of depression than work-derived stress (5). People who live an involuntary solitary life tend to have more suicidal ideation than others in the general population (6). Several studies have reported that the recent COVID-19 pandemic–related social isolation has increased the prevalence of depression and anxiety (7, 8).

A plausible mechanism for depression is known as the serotonin depletion hypothesis (9). Thus, medications targeting the serotonergic system, such as selective serotonin reuptake inhibitors (SSRIs), are the most frequently prescribed medications to patients with depressive disorders (10). However, these antidepressants have limitations such as a low response rate (approximately 47%), a high remission rate (approximately 28%), and a high risk of relapse following drug withdrawal (approximately 62% with fluoxetine) (11–13). Recently, neuroinflammation has been proposed as one pathophysiologic basis of depression and is considered a promising therapeutic target (14, 15).

On the other hand, to escape from sadness and melancholy, individuals with depression are prone to addiction, especially excessive alcohol use (16). Clinical reports have shown that approximately 40.3% of patients with major depressive disorder (MDD) have alcohol use disorder (AUD) (17), and conversely, 63.8% of those with AUD have a high prevalence of major depression (18). Alcohol use magnifies suicide risk and leads to serotonergic hypoactivity in patients with depressive disorder (19). As expected, uncontrolled alcohol use is known to promote the pathological progression and to hinder the treatment of depression (20, 21). Previously, the transcriptome analysis showed increased levels of translocator protein in activated microglia in the human alcoholic brain as a neuroinflammatory feature (22). However, the pathological mechanisms of the negative mutual interaction between depression and alcohol consumption remain unclear. Here, we used a mouse model involving social isolation stress (IS) and/or voluntary ethanol intake and explored the mechanisms underlying the interaction between IS-induced depression and ethanol-seeking behaviors.

RESULTS
Voluntary ethanol intake during IS
IS increased voluntary ethanol intake (but not tap water intake) approximately 2.0–2.5-fold over the course of 28 days compared to that in the control group (P < 0.05 and P < 0.01; Fig. 1, A and B). This phenomenon tended to be similar in female mice (fig. S2D). Meanwhile, restraint stress partially increased ethanol intake (P < 0.01; Fig. 1C). Sucrose preference was significantly reduced by restraint stress (P < 0.05), but it was not reduced by IS (Fig. 1D).

Ethanol CPP during IS
From the conditioned place preference (CPP) test, an increased ethanol place preference was observed in mice subjected to IS. The ethanol place preference score in the reinstatement test following extinction (for 8 days) was significantly elevated in the IS group compared to both the control and saline-injected groups (P < 0.01 and P < 0.05, respectively; fig. S2E).

Changes in body weight and food intake
The mice gained approximately 2 g of body weight after 4 weeks, and changes in body weight among the groups were not affected by IS and/or ethanol intake (fig. S2A). During the experimental days, food intake tended to decrease in mice subjected to IS along with ethanol but was not significantly different among the four groups (fig. S2B).
Changes in serum ethanol concentrations and serum levels of aspartate transaminase and alanine transaminase

Serum ethanol concentrations were significantly increased by ethanol intake, irrespective of whether mice were subjected to IS (P < 0.05 for both) or not (fig. S2G). Serum aspartate transaminase (AST) and alanine transaminase (ALT) levels were not altered by IS and/or voluntary ethanol intake (fig. S2, H and I).

Changes in anxiety- and depressive-like behaviors

In the open field test (OFT; total distance traveled and time spent in the center) and elevated plus maze (EPM; distance traveled, number of arm entries, number of open arm entries, and time spent time in the open arms), IS significantly induced anxiety-like behaviors (P < 0.05 or P < 0.01) compared to those in the respective control groups. These alterations following IS were remarkably exacerbated by ethanol intake (P < 0.05 and P = 0.071; Fig. 2A, a and b; P < 0.05 for all parameters; Fig. 2B, a to d).

The results from both the forced swimming test (FST; total global activity, immobility duration, and activity duration) and tail suspension test (TST; total global activity, immobility duration, and latency to immobility) indicated that IS induced a considerable increase in depressive-like behaviors (P < 0.05 or P < 0.01) compared to those in the respective control groups. In particular, depressive-like behaviors were most intense in mice that both were exposed to IS and had access to ethanol (P < 0.05 and P = 0.081, respectively; Fig. 2C, a and b; P < 0.05 for all parameters; Fig. 2D, a to c). In contrast, the mice receiving a single injection of ethanol (1 g/kg, intraperitoneally) at night (02:00 hours, main active hour) showed a slight alleviation of depressive-like behaviors, as shown in the FST 12 hours later at 14:00 hours (P < 0.05; fig. S2C).

Changes in serotonergic signals and microglial activity

Serotonergic activity in the raphe nuclei (dorsal and median) was prominently inhibited by IS, as shown by both 5-hydroxytryptamine (5-HT; P < 0.01) and tryptophan hydroxylase 2 (TPH2; P < 0.01) staining, compared to that in control conditions (Fig. 3 and fig. S3) but not by ethanol intake alone. These two suppressed signals were further emphasized in the dorsal raphe nuclei of mice subjected to IS along with ethanol (P < 0.05 for both; Fig. 3, A and B, and fig. S3, A to D).

In contrast to serotonergic activity, activation of microglia, as evaluated by increases in the intensity, numbers, and soma area of the Iba-1–positive cells in the dorsal raphe nucleus, was greater in the IS group than in the respective control groups (P < 0.05 or P < 0.01; Fig. 3, A and C to E). Moreover, microglial activation was further increased by ethanol intake (P < 0.05 or P < 0.01; Fig. 3, A and C to E). This phenomenon was also observed by immunohistochemical staining with Iba-1 (P < 0.01; fig. S3, E and F).

Changes in microglial function and inflammatory cytokine

IS significantly increased microglial activation (Iba-1) and phagocytosis-derived molecules involved in synaptic pruning (C1q and cleaved...
Fig. 2. Depressive- and anxiety-like behaviors. After IS with or without ethanol exposure, the total distance traveled (a) and time spent in the center (b) in the OFT (A) were assessed on the 28th day. After a period of 6 hours, total global activity (a), immobility duration (b), and activity duration (c) in the TST (C) were assessed. The traveled distance (a), total number of entries (b), entries into the open arms (c), and spent time in the open arms (d) in the EPM (B) were assessed on the 29th day. Total global activity (a), immobility duration (b), and latency to immobility (c) in the FST (D) were assessed on the 30th day. The data are expressed as the means ± SD. *(a) P < 0.05 compared to the unstressed mice, and *(b) P < 0.05 compared to the unstressed mice.

Accordingly, microglia-mediated neuroinflammation was triggered by IS in the raphe nuclei, as evidenced by differences in the levels of proinflammatory cytokines (tumor necrosis factor, TNF-α) and anti-inflammatory cytokines (interleukin, IL-10) compared to those in the respective control groups (P < 0.05 for both parameters; Fig. 4, A and B). These neuroinflammatory responses were noticeably further enhanced by exposure to ethanol intake (P < 0.01 for both parameters; Fig. 4, C and D). To confirm whether ethanol directly activates microglia, BV2 cells were activated by lipopolysaccharide (LPS). The gene expression levels of TNF-α and IL-1β were significantly increased by ethanol treatment (150 mM; P < 0.05 and P < 0.01, respectively; Fig. 4, E and F). The low dose of ethanol (20 mM) also increased IL-1β mRNA levels (P < 0.05; fig. S2F).

**Effects of a microglial inhibitor**

Administration of minocycline (a selective microglial inhibitor) completely prevented microglial-mediated loss of the serotonergic indicators (Fig. 5, A and B). Minocycline also alleviated the IS-induced overactivation of microglia (Fig. 5, C and D), which was supported by differences in C1q protein and C1qa mRNA levels (P < 0.05;
Fig. 5E). Moreover, minocycline significantly improved depressive-like behaviors, as shown in the FST, including immobility duration and latency to immobility ($P < 0.05$ and $P < 0.01$, respectively; Fig. 5, F and G).

**Changes in the mesolimbic dopaminergic system**
In the mice exposed to a socially isolated environment, there were increases in both tyrosine hydroxylase (TH; $P < 0.01$) and dopamine ($P < 0.01$) levels in the ventral tegmental area (VTA). These two signals were markedly higher in mice subjected to IS and ethanol than in mice subjected to stress only ($P < 0.05$ and $P < 0.01$, respectively; Fig. 6, A to C). Significant increases in the dopaminergic projections into the nucleus accumbens (NAc) were shown by c-Fos translocation to the nucleus (fig. S4A).

**DISCUSSION**
The above results clearly demonstrated the mutual interaction between depressive-like behaviors and ethanol consumption. Although free access was given to both water and ethanol (the two-bottle choice procedure), social IS markedly induced a radical increase in ethanol intake (i.e., sixfold increase at approximately 2 weeks; Fig. 1A and movies S1 and S2). To verify ethanol-seeking behavior, we additionally assessed ethanol-induced CPP using a tactile conditioned stimulus. The ethanol preference score (on the day of the reinstatement test) was considerably increased by IS (fig. S2E). We confirmed that ethanol consumption was much more prominent during IS conditions than in conditions involving restraint stress (Fig. 1C). The serum ethanol concentrations were significantly increased in the two groups given the choice of ethanol intake. However, there was no significant difference between the IS and the control group (assessed at approximately 14:00 hours) in blood ethanol concentration (fig. S2G), which might be explained by the rapid metabolism; ethanol in the C57BL/6J mouse is metabolized within 5 hours in the blood ($23$). Among the groups, metabolism rates may not be different because the liver was intact (fig. S2, H and I). An additional experiment confirmed that ethanol overconsumption was exhibited by female mice (fig. S2D).

Notably, IS-induced depressive-like and anxiety-related behaviors were aggravated by ethanol consumption, as we expected...
significantly exacerbated by ethanol intake (Fig. 3 and fig. S3). The
in the raphe nuclei. Moreover, this hyposerotonergic activity was
social IS conspicuously reduced 5-HT (serotonin) and TPH2 signals
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Fig. 4. Microglia synaptic pruning and neuroinflammation. After IS with or without ethanol exposure for 28 days, mice were euthanized. Raphe nuclei homogenates were used to analyze protein and cytokine expression. The protein levels of Iba-1, PSD95, synaptophysin, cleaved caspase 3, and C1q were determined by Western blotting (A), and their intensities were semiquantified (B). Levels of the cytokines TNF-α (C) and IL-10 (D) in the raphe nuclei were determined by enzyme immunoassay. The data are expressed as the means ± SD (n = 5). *P < 0.05 and **P < 0.01 compared to the vehicle-treated cells. 

(Fig. 2, A to D). We verified that these behavioral changes were not induced by acute ethanol exposure (fig. S2C). Previous animal studies using IS models, such as maternal separation or adolescent social isolation, have shown a similar pattern of ethanol craving (24–26). Poor relationships with friends or family members are considered to be a factor underlying the co-occurrence of both depression and AUD (27), and their comorbidity is a significant concern worldwide (28). Heavy or frequent drinking increases the risk of depressive symptoms during adolescence (twofold higher than in adulthood) (29). Under the current COVID-19 pandemic crisis condition, clinical studies have revealed a positive association between social isolation and increases in depressive disorder and alcohol misuse (7, 30, 31). However, the mechanisms underlying this interaction remain unclear.

To explain the interactive mechanisms, we examined relevant neurotransmitter systems and related mediators. Sociality is regulated by serotonergic innervation from the raphe nuclei (32, 33), and it plays a pivotal role in coping with inescapable stress (34). Serotonin depletion is a common feature of the pathophysiology of MDD (35, 36), and a previous meta-analysis found a functional genetic variant of TPH2 (a rate-limiting enzyme in the synthesis of 5-HT) in patients with major depression (37). As we expected, social IS conspicuously reduced 5-HT (serotonin) and TPH2 signals in the raphe nuclei. Moreover, this hyposerotonergic activity was significantly exacerbated by ethanol intake (Fig. 3 and fig. S3). The alcohol-mediated disruption of the serotonergic system has been indicated in both clinical- and animal-based data: 5-HT depletion has been observed in the cerebrospinal fluid of those with AUD and in the raphe nuclei of adolescent rats exposed to ethanol binge (38, 39). We thus speculate that social isolation–derived serotonergic depletion causes depressive-like behaviors and sequentially leads to ethanol-seeking and vice versa.

Drinking habits are an obstacle in the course of treatment for depressive disorders in clinics, which leads to a reduced response, up to 70%, to SSRIs (40). Recently, microglia have emerged as a promising target for the treatment of depression (41, 42). A positron emission tomography (PET) imaging study found microglial activation in patients with depression (43). An animal study evidenced microglial hyperactivity lasting for more than 30 days after 4-day binge alcohol exposure (44). In our results, concomitant IS and ethanol intake activated microglia around serotonergic neuronal cells in the dorsal raphe nucleus in an additive manner (Fig. 3). These results might be supported by other data showing the connection between microglial activation and 5-HT neuronal inactivity (45, 46). Microglia have emerged as important players in the synapse pruning process in both developing and aging brains (47–49), and their by-product C1q, microglial-derived complement cascade protein, induces neuronal cell death (50). Positive correlations between high serum C1q levels and the severity of depression were observed in patients with MDD (51). As expected, IS and ethanol
consumption led to considerable synaptic loss, as evidenced by the increased levels of Iba-1, C1q, and cleaved caspase 3, but decreases in synaptophysin and PSD95 (indicators of pre- and postsynaptic activity, respectively) in the raphe nuclei (Fig. 4, A and B).

Regarding the involvement of microglia in alcohol-related behaviors and molecular alterations, we conducted a confirmatory experiment using minocycline, a selective inhibitor of microglial activation. Minocycline has pharmacological properties against several neurological disorders by inhibiting microglial activation and neuroinflammation–related molecules, such as caspase-1–dependent IL-1β and inducible nitric oxide (52). As expected, treatment with minocycline significantly attenuated microglial hyperactivation, synaptic pruning, and depressive-like behaviors (Fig. 5). A double-blind randomized clinical trial revealed the therapeutic potential of minocycline in patients with MDD tolerant to commonly used antidepressant (53). These results indicated the pathogenetic roles of microglial hyperactivation in serotonergic damage in IS and ethanol overconsumption conditions. On the other hand, scientists well

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Fig. 5. Effects of minocycline on serotonergic depletion, microglial activation, ethanol intake, and depressive-like behavior. Following IS with ethanol exposure and minocycline injection for 28 days, the mice were euthanized. Microglial-derived serotonergic depletion was confirmed by immunofluorescence analysis of 5-HT and Iba-1 (A), and microglial activity was assessed by costaining of Iba-1 and C1q (C) in the dorsal raphe nuclei. Simple linear regression analyses were performed (B and D). Raphe nuclei homogenates were used to analyze C1q gene expression (E). In another experiment, ethanol intake (F) was assessed, and depressive-like behavior was assessed using the FST (G). The data are expressed as the means ± SD (n = 3 or 8). **P < 0.01 compared to the unstressed mice, and #P < 0.05 and ##P < 0.01 compared to the mice not injected with minocycline. DAPI, 4′,6-diamidino-2-phenylindole; GAPDH, glyceraldehyde-3-phosphate dehydrogenase.
noticed the vital functions of activated microglia-derived proinflammatory cytokines in the poor sociality and depressive-like behavior (54, 55). Our data revealed the IS-derived alterations in TNF-α and IL-10 levels in the brain and exacerbation by exposure to ethanol (Fig. 4, C and D). Ethanol directly increased TNF-α and IL-1β gene expression in a BV2 microglial cell line (Fig. 4, E and F).

We further investigated the mechanisms underlying ethanol craving from the perspective of the reward system. Dysregulated reward circuitry is common among patients with substance (nicotine and alcohol) use disorders and neuropsychiatric disorders including depression or anxiety (56). Compared to artificial stimuli such as restraint or foot shock stress, social defeat–like stress is known to sensitize rodents to ethanol-or cocaine-related rewards (57). In our study, the mice that were exposed to restraint stress did not exhibit a strong response to ethanol (Fig. 1C), while IS markedly increased both TH and dopamine (Fig. 6, A to C). Moreover, IS produced changes in neuronal firing (c-Fos–positive activity) in NAc, although changes in the dopaminergic project from VTA were uncertain (fig. S4). One group observed higher activity of dopaminergic reward circuits in patients with social anxiety disorder than in healthy controls using functional magnetic resonance imaging analysis (58).

Although we conducted a comprehensive experiment, there are some limitations, for example, adoption of only a single animal model and one age group (6-week-old mice). Compared with adult rodents, young rodents are known to be more susceptible to voluntary ethanol consumption during IS conditions (51). Although each variable (social isolation, ethanol, and neuroinflammation) is known as a risk factor, the novel findings from this study are as follows: (i) Our findings provide the first evidence to verify the interconnections among these factors, especially regarding the vicious cycle of the relationship; (ii) we are the first to provide these novel mechanistic insights into the causative role of microglia in clinical comorbidity between depressive disorders and AUDs, as summarized in fig. S5; and (iii) these are interesting findings that verify the mutual specificity of IS (not restraint stress) and ethanol (not sucrose), respectively. Our findings provide insight and ideas to physicians and scientists to develop a therapeutic strategy for the growing number of depressed patients living in solitary environments and their alcohol issues.

MATERIALS AND METHODS
Animal and experimental design
Specific pathogen–free male adolescent C57BL/6J mice (5 weeks old, 21 to 23 g) were purchased from Dae Han Bio Link Co. Ltd. (Eumseong, Republic of Korea). They were maintained under a thermo hygrostat (temperature: 23°C ± 2°C and humidity: 55 ± 10%; ALFFIZ, Busung Co. Ltd., Seoul, Republic of Korea) on a 12-hour light-dark cycle (09:00 to 21:00 hours), freely fed food pellets (Cargill Agri Purina, Gyeonggi-do, Republic of Korea), and provided two bottles of tap water before the experiments began. For all experiments, the mice, except those in the control groups (three to five mice per cage), were individually housed for 28 days to induce a depressive-like state. Voluntary ethanol intake, food intake, and body weight were recorded over these 28 days (twice per week).

Three main studies were conducted as described below. After acclimation for 1 week, the mice were randomly divided into the following three experiments (n = 32 per experiment): Experiment 1,
which was performed to assess ethanol-related behaviors and depressive- and anxiety-like behaviors; experiment 2, which was performed to analyze cytokine and immunohistological alterations; and experiment 3, which was performed to determine serum biochemical and protein alterations. The mice in each experiment were divided into the following four groups ($n = 8$ per group) based on two variables, namely, social IS and voluntary ethanol intake: control with two bottles of tap water, control with both tap water and 10% ethanol, IS with two bottles of tap water, and IS with both tap water and 10% ethanol. During the 1-week acclimation period, the mice were adapted to the two-bottle choice procedure (two bottles containing tap water). To achieve a human-like approach to drinking, mice were allowed to spontaneously consume tap water and/or 10% ethanol for 28 days, with the positioning of the two bottles being changed at 4-day intervals.

To conduct six additional studies, the experiments were designed as follows: experiment 4, which was performed to evaluate the influence of minocycline ($n = 3$ or $8$ per group)—control group (tap water and 10% ethanol), IS group (tap water and 10% ethanol), and IS group (tap water and 10% ethanol) receiving minocycline (50 mg/kg, intraperitoneally; daily for 28 days); experiment 5, which was performed to verify specificity of IS effects ($n = 5$ per group)—control group (tap water and 10% ethanol) and group (tap water and 10% ethanol) exposed to restraint stress as previously described (3 hours per day for 28 days); experiment 6, which was performed to determine specificity of ethanol effects ($n = 5$ per group)—control group (tap water and 10% ethanol) and group (tap water and 10% ethanol and 1% sucrose) and social IS group (tap water and 1% sucrose); experiment 7, which was performed in female mice ($n = 3$ per group)—control group (tap water and 10% ethanol) and IS group (tap water and 10% ethanol); experiment 8, which was performed to confirm ethanol-seeking behaviors ($n = 10$ per group); and experiment 9, which was to investigate acute ethanol effects on behavioral alterations ($n = 6$ per group). All experiments were repeated two to three times with the indicated numbers of mice. The experimental designs are displayed in fig. S1. All of the experiments were approved by the Institutional Animal Care and Use Committee of Daejeon University (DJUARB2016-037) and were conducted in accordance with the Guide for the Care and Use of Laboratory Animals, published by the National Institutes of Health (NIH).

**Social IS protocol**
The social IS procedure was performed as described previously (59). Briefly, the mice were individually housed in a single cage with a screen blocking visual communication for 28 days. During the period of isolation, the mice were exposed to minimal handling and were provided no environmental enrichment.

**Two-bottle drinking procedure**
The mice were subjected to a two-bottle choice procedure as previously described (60). The mice were given continuous access to either two bottles of tap water or a bottle of tap water and a bottle of 10% ethanol (v/v) in each experiment. The positioning of the two bottles (left and right) was changed at 4-day intervals to avoid the effect of preferences for a particular side, and intake was recorded.

**Behavioral monitoring (experiment 1) Open field test**
Anxiety-like behavior of the mice was first tested in a large square chamber (30-cm height × 40-cm length × 40-cm width). To assess activity, the open field arena was equally divided into quadrants (20 × 20 cm), and a video camera was mounted overhead. The mice were allowed to freely explore the test arena for 5 min. After each test, the apparatus was cleaned with acetic acid (2%) to limit mouse odors. The total horizontal distance (meters) and the time spent in the center (seconds) were recorded at 50 lux illuminations on the 28th day.

**Tail suspension test**
Depressive-like behavior was measured using a rectangular compartment (55-cm height × 15-cm width × 12-cm depth) with an aluminum suspension bar (1 × 1 cm; positioned on the top of the box). Mouse tails were fastened with tape to the suspension bar, and the suspended mice were observed for 5 min. Level of immobility was determined on the basis of the designated thresholds for immobility duration (immobile: 0 to 120, active: 121 to 300). Total global activity, immobility, and activity durations (seconds) were recorded at 25 lux illuminations on the 28th day.

**Elevated plus maze test**
The mice were subjected to another assessment of anxiety-like behavior. The apparatus consisted of two opposing open arms (30-cm arm length × 5-cm arm width at 20 lux) and two closed arms (30-cm arm length × 5-cm arm width × 15-cm high walls at 90 lux) extending from a central area (5 × 5 cm) and elevated above the ground (20 cm). Each mouse was placed in the central area of the maze with access to all arms and was allowed to freely explore the maze for 5 min. The total distance traveled (centimeters), number of entries into the open arms, and time spent in the open arms were recorded on the 29th day.

**Forced swimming test**
The mice were subjected to the FST to measure despair-related behavior. The apparatus consisted of a plastic cylinder (30-cm height × 20-cm diameter) filled with water to 10 cm from the top at 24° ± 1°C. Each mouse was allowed to freely swim for 5 min. Level of immobility was determined on the basis of the designated thresholds for immobility duration (immobile: 0 to 120, active: 121 to 300) established by the Smart global activity module. Total global activity, immobility duration, and latency to immobility (seconds) were recorded at 30 lux illuminations on the 30th day.

**Conditioned place preference**
To confirm ethanol-seeking behaviors, the CPP test was adopted. The apparatus (MED-CPP-3013-2, Med Associates, VT, USA) consisted of two compartments (17.4-cm height × 12.7-cm length × 12.7-cm width at 10 lux) with a guillotine door separating the compartments. Infrared beams detected the movement of each mouse in the black (grid floor) and white (hole floor) compartments. During the preconditioning test, mice exhibiting an unconditioned preference (more than 800 s on either side) for either compartment were excluded. The ethanol-CPP paradigm was performed as follows: (i) The mice were allowed to freely explore the compartments for 20 min (preconditioning test on the first day). From the next day to the ninth day after preconditioning habituation, each mouse was confined to the respective compartment with discriminable tactile cues (grid or hole floor) for 5 min following 10 min after injection of saline or ethanol (2 g/kg, intraperitoneally). (ii) The time each mouse spent in each compartments was recorded for 20 min (postconditioning test at 10th day). (iii) On the seventh day after extinction session, the mice were tested for 20 min immediately following saline or ethanol (1 g/kg, intraperitoneally) injection (reinstatement test at 18th day). The following formula was used to calculate the
The day after the last day of IS, the mice were euthanized under CO₂ anesthesia conducted in the daytime (13:00 to 17:00 hours). Serum was collected from abdominal blood by centrifugation at 3000g for 15 min. The brains of five mice from each group were immediately removed. Brain regions were isolated using a coronal mouse brain matrix (1 mm; BSMAS001-1, Zivic Instruments) and biopsy punch (1 mm; BP-10F, Kai Medical). For biochemical analysis, samples were homogenized in radioimmunoprecipitation assay buffer (R0278, Sigma-Alrich) supplemented with protease inhibitors (no. 1183615001, Roche). The total protein concentration was measured using a bicinchoninic acid protein assay kit (BCA1 and B9643, Sigma-Alrich). The absorbance was measured using an ultraviolet (UV) spectrophotometer at 562 nm (Molecular Devices Corp., Sunnyvale, CA, USA). For immunohistochemical analyses, the three remaining mice from each group were transcardially perfused with 0.05% heparin (10 U/ml in phosphate-buffered saline (PBS)) followed by 4% paraformaldehyde (pH 6.9). The brains were placed in the same solution for fixation.

**Immunohistochemical analysis**

Immunohistochemical staining was performed to assess microglial activity (Iba-1), serotonergic activity (5-HT and TH2PH) in the raphe nuclei, dopaminergic activity (TH and dopamine) in the VTA, and neural activity (c-Fos) in the NAc. The brains were gradually cryo-protected in 10, 20, and 30% sucrose for 24 hours each and were subsequently embedded in optimal cutting temperature compound (CM3050_S, Leica) in liquid nitrogen. They were cut into frozen coronal sections (35 μm) using a cryostat (CM3050_S, Leica). The sections were stored free floating in buffer. For washing with ice-cold PBS, parallel free-floating sections were treated with blocking buffer (5% normal chicken serum in PBS and 0.3% Triton X-100 for 1 hour at 4°C) and incubated with anti-rabbit Iba-1 monoclonal (1:400; no. 019-19741, Wako Biologicals), anti-rat C1q monoclonal (1:10; ab11861, Abcam), anti-goat 5-HT monoclonal (1:200; ab66047, Abcam), anti-rabbit TH2polyclonal (1:500; NB100-74555, Novus), anti-rabbit TH polyclonal (1:500; NB300-109, Novus), anti-mouse dopamine monoclonal (1:500; NB110-2538, Novus), and anti-rabbit c-Fos polyclonal (1:100; NBP1-74555, Novus) primary antibodies overnight at 4°C. After washing with ice-cold PBS, the sections were incubated with a donkey anti-goat immunoglobulin G (IgG) H&L (1:400; Alexa Fluor 488, ab150129), goat anti-rabbit IgG H&L (1:400; Alexa Fluor 488, ab150077), goat anti-mouse IgG H&L (1:400; Alexa Fluor 488, ab150113), goat anti-rabbit IgG H&L (1:400; Alexa Fluor 594, ab150080), goat anti-mouse IgG H&L (1:400; Alexa Fluor 594, ab150160), or goat anti-rat (1:400; Alexa Fluor 594, ab150160) secondary antibodies for 2 hours at 4°C. The sections were subsequently exposed to 4',6-diamidino-2-phenylindole (1:1000; D9542, Sigma-Alrich) to stain the cell nuclei.

Immunohistochemical analysis of Iba-1 (microglial activity) was performed in the dorsal raphe nucleus. The sections were incubated with a primary antibody against Iba-1 (1:200; no. 019-19741, Wako Biologicals) overnight at 4°C. The sections were incubated with a horseradish peroxidase (HRP)–conjugated goat anti-rabbit IgG (1:400; ab6722, Abcam) secondary antibody for 2 hours at room temperature. For amplification of the signal, the sections were exposed to an avidin–biotin peroxidase complex (Vectastain ABC kit, Vector Laboratories) for 2 hours. Peroxidase activity was visualized using stable diaminobenzidine solution. Immunoreactivity was observed under an AxioPhot microscope (Carl Zeiss, Germany). Signals were quantified using ImageJ 1.46 software (NIH, Bethesda, MD, USA), and morphological characteristics of stained microglia (average cell body size per cell, average dendritic process per cell, and cell number per mm²) were analyzed by using image analysis software (Image-Pro Plus 6.0, Media Cybernetics Inc., Rockville, USA).

**Western blot analysis**

The protein expression levels of Iba-1, C1q, cleaved caspase 3, synaptophysin, PSD95, and β-actin in raphe nuclei homogenates were evaluated using Western blotting. The protein concentrations of the homogenates were equalized, and the samples were separated by 10% polyacrylamide gel electrophoresis and transferred to polyvinylidene fluoride membranes. To minimize nonspecific binding, the membranes were blocked in 5% bovine serum albumin for 1 hour. The membranes were incubated overnight at 4°C with the following primary antibodies: Iba-1 (1:500; 061-20001, Wako Biologicals), C1q (1:50; ab71940, Abcam), cleaved caspase 3 (1:200; no. 9664S, Cell Signaling Technology), PSD95 (1:200; ab13552, Abcam), synaptophysin (1:500; ab8049, Abcam), or β-actin (1:2500; PA1-183, Thermo Fisher Scientific). After washing, the membranes were incubated with an HRP-conjugated anti-rabbit or anti-mouse antibody (GeneTex Inc., Irvine, CA) for 1 hour. The Western blotting results were visualized with an enhanced chemiluminescence advanced kit. The intensity was analyzed by ImageJ version 1.46 (NIH, Bethesda, MD, USA).

**Pro- and anti-inflammatory cytokines**

TNF-α and IL-10 levels in raphe nuclei homogenates were determined using commercially available enzyme immunoassay (OptEIA) kits for TNF-α (no. 558534, BD Biosciences, San Diego, CA, USA) and IL-10 (no. 555252, BD Biosciences, San Diego, CA, USA), and the absorbance was measured at 450 and 570 nm using a UV spectrophotometer (Molecular Devices Corp., Sunnyvale, CA, USA).

**Ethanol concentrations**

The serum levels of ethanol (K610, BioVision, Milpitas, CA, USA) were measured using an ethanol colorimetric assay kit. Absorbance was measured at 450 or 570 nm using a UV spectrophotometer (Molecular Devices Corp., Sunnyvale, CA, USA).

**AST and ALT levels**

The serum levels of AST and ALT were determined using an autoanalyzer (Chiron Diagnostics Co.).

**Real-time quantitative polymerase chain reaction (in vitro experiment)**

Mouse microglial cells (BV2 cells) were cultured in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum and 1% penicillin-streptomycin. BV2 cells were incubated at
37°C under 5% CO₂. BV2 cells were seeded into six-well plates at a density of 2 × 10⁵ cells per well. After incubation for 12 hours, the BV2 cells were treated with PBS or LPS (100 ng/ml) from Escherichia coli O111:B4 (no. L2630, Sigma-Aldrich) or 10, 20, or 150 mM ethanol for 24 hours. Ethanol-treated cells were incubated under a beaker containing 200 ml of 4% ethanol to prevent the evaporation of ethanol from the media. To confirm whether ethanol induces microglial activation, the gene expression of inflammatory cytokines was measured in cell lysates.

The mRNA expression of genes encoding TNF-α, IL-1β, and glyceraldehyde-3-phosphate dehydrogenase in BV2 cells or C1q in raphé nuclei homogenates from mice was measured by real-time polymerase chain reaction (PCR). Total mRNA was extracted using an RNeasy Mini Kit (QIAGEN, Valencia, CA, USA), and cDNA was synthesized using a High-Capacity cDNA Reverse Transcription Kit (Ambion, Austin, TX, USA). Real-time PCR was performed using SYBR Green PCR Master Mix (Applied Biosystems, Foster City, CA, USA), and PCR amplification was performed using a standard protocol on an IQ5 PCR Thermal Cycler (Bio-Rad, Hercules, CA, USA). Information regarding the primer sequences is summarized in table S1.

Statistical analysis
All results are expressed as the means ± SD. Statistical analysis was performed by using two-way analysis of variance (ANOVA) followed by post hoc analysis by Bonferroni t test with exposure (control or IS) and drink (tap water or ethanol) as the between-subject variables using IBM Statistical Package for the Social Sciences (SPSS) statistics software, ver. 25.0 (SPSS Inc., Chicago, IL, USA). Differences at P < 0.05 indicate statistical significance. For correlations, a linear regression analysis was performed.

SUPPLEMENTARY MATERIALS
Supplementary material for this article is available at https://science.org/doi/10.1126/sciadv.ajb3400

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