Repeated Sevoflurane Exposures in Neonatal Rats Increased the Brain Vulnerability to Future Stress Exposure and Resulted in Fear Extinction Deficit

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Abstract
Sevoflurane anesthesia during neonatal period was reported to sensitize the rodent animals to stress later in life. The authors tested the hypothesis that repeated sevoflurane exposures in neonatal rats increased the brain vulnerability to future stress exposure and resulted in fear extinction deficit and investigated whether the neonatal brain depolarizing γ-aminobutyric acid type A receptor (GABA\(_A\)R) is involved in mediating these abnormalities. Neonatal Sprague–Dawley male rats, pretreated with vehicle or the NKCC1 inhibitor, bumetanide, received sequential exposures to 3% sevoflurane for 2 h on postnatal days (P) 5, P6, and P7 and then were exposed to electric foot shock stress in fear conditioning training at P14. Juvenile rats at different developmental brain stage receiving identical sevoflurane exposures on P25, P26, and P27 were also studied. The results showed repeated sevoflurane exposures in neonatal rats and increased the cation-chloride cotransporters NKCC1/KCC2 ratio in the PFC at P14. Repeated exposures to sevoflurane in neonatal rather than juvenile rats enhanced the stress response and exacerbated neuroapoptosis in the PFC after exposed to electric foot shock in fear conditioning training. Neonatal rather than juvenile sevoflurane-exposed rats exhibited deficits in fear extinction training and recall. Pretreatment of neonatal rats prior to sevoflurane exposures with bumetanide reduced the NKCC1/KCC2 ratio at P14 and ameliorated most of the subsequent adverse effects. Our study indicates that repeated sevoflurane exposures in neonatal rats might increase the brain vulnerability to future stress exposure and resulted in fear extinction deficit, which might be associated with the neonatal enhanced brain depolarizing GABA\(_A\)R activity.

Keywords Neonatal sevoflurane · Stress exposure · Neuroapoptosis · Fear extinction · NKCC1/KCC2 ratio

Introduction
Sevoflurane is routinely administered to infant of humans for procedural anesthesia. Neonatal sevoflurane-induced neurobehavioral dysfunctions have been displayed in some animal experimental reports (Yu et al. 2020, 2018; Ji et al. 2017; Sun et al. 2016). However, since there are various environmental challenges or stress during brain development for humans, the association between neonatal sevoflurane exposure and future neurocognitive disorders were always full of controversy.

Disturbance of neuroendocrine response to stress were reported to be associated with many neuropsychiatric disorders, such as depression and posttraumatic stress disorder (PTSD) (Ménard et al. 2017; Molai and Maguire 2018; DePierro et al. 2019). People who used to be suffered from extremely stress events have an increased risk in development of PTSD. In recent years, more and more animal
experimental reports have demonstrated that early life anesthetic exposure might contribute to neuroendocrine disturbance and sensitize the animal to environmental stress later in life (Tan et al. 2014; Xu et al. 2015; Ju et al. 2017). Moreover, later life environmental insult or stress might exacerbate neurodevelopmental disorders for neonatal anesthetically exposed animals (Ju et al. 2017). It has been considered that the adverse developmental results of anesthetic exposure in neonatal animals might be linked with future experiences (Zhang et al. 2015); it is plausible that neurocognitive abnormalities might be the result of an accumulated influence initially programmed by early life anesthetic exposure and later exacerbated by future adverse stressful factors, such as various types of psychic and physical trauma.

Sevoflurane exposure at early life for rodent animals might contribute to abnormal social behaviors similar to autism spectrum disorder (ASD) and neuropsychiatric-like behavior changes such as social interaction deficits (Satomo et al. 2009; Lin et al. 2016). It has been reported that certain traits of ASD might predispose the population to an increased vulnerability to future adverse stress exposure and development of PTSD (Kerns et al. 2015; Hoover 2015; Haruvi-Lamdan et al. 2018). The activation targets of sevoflurane include γ-aminobutyric acid (GABA) type A receptor (GABA_A R).

Early-life sevoflurane exposure impaired GABA_A R-mediated excitatory/inhibitory balance and enhanced seizure activity later in life (Lin et al. 2021). The cation-chloride cotransporters Na^+-K^+-2Cl^− (NKCC1) and K^+-2Cl^−-2 (KCC2) play a crucial role in regulation of GABA_A R response to GABA. NKCC1 renders GABA_A R excitatory to GABA in the immature neurons, whereas the increased expression of KCC2 renders GABA_A R to its inhibitory role. Imbalances in transition of NKCC1/KCC2 could lead to hyperexcitability in neurocircuitry. The NKCC1 inhibitor bumetanide was reported to enhance the role of GABAergic inhibition, which attenuated the electrical and behavioral disorders in models of ASD (Lemonnier et al. 2017; Tzyio et al. 2014; Dzhala et al. 2005).

In the present study, the neonatal sevoflurane-exposed rats were exposed to electric foot shock stress in fear conditioning training, and the authors tested the hypothesis that repeated sevoflurane exposures in neonatal rats increased the brain depolarizing GABA_A R is involved in mediating these abnormalities.

### Material and Methods

#### Animals

The present study was approved by the Ethics Committee of West China Hospital, Sichuan University, and was guided in accordance with the ARRIVE guidelines and the Guide for the Care and Use of Laboratory Animals from the National Institutes of Health (Bethesda, MD, USA). Pregnant Sprague–Dawley rats were housed individually in standard conditions with a 12 h light/dark cycle (light from 07:00–19:00) at 24 ± 1 °C and ad libitum access to food and water. Five days after the delivery, litters were culled to 12 male pups. Pups were weaned at P21 and housed in groups of four for a further study during juvenile period. To control for litter variability, we used several male pups from different litters for each treatment condition.

#### Experimental Design of the Study

The experimental design of the study is illustrated in Fig. 1. In the first set of experiments (Fig. 1A), the neonatal P5 to P7 male rat pups were randomly assigned to sevoflurane (SEV) and control (CON) groups, and then rats were exposed to electric foot shock stress in fear conditioning training at P14. The NKCC1/KCC2 ratio in the prefrontal cortex (PFC) was determined after rats exposed to electric foot shock stress in fear conditioning training. To test the stress response and brain vulnerability to future stress exposure after repeated sevoflurane exposures in neonatal rats, serum corticosterone and neuroapoptosis in the PFC after exposed to electric foot shock stress in fear conditioning training at P14 were measured. To test the hypothesis that repeated sevoflurane exposures in neonatal rats resulted in future fear extinction deficit, fear conditioned rats were subjected to fear extinction training at P18. Fear extinction recall was assessed at P20. In the second set of experiments (Fig. 1B), in order to test whether sevoflurane exposure at different developmental brain stage contributed to different

![Fig. 1 Experimental design of the study](image-url)
results, juvenile rats receiving identical sevoflurane exposures on P25, P26, and P27 were explored.

**Neonatal and Juvenile Sevoflurane Exposure**

The neonatal P5 to P7 male rat pups were randomly assigned to sevoflurane (SEV) or control (CON, not exposed to sevoflurane) groups. Rats in the SEV group were exposed to 3% sevoflurane in O2/N2 (fraction of inspired oxygen 50%, or FiO2 50%) for 2 h daily on 3 consecutive days from P5 to P7, in a thermostated chamber set to 37 ± 1 °C. The total gas flow was two Litre min⁻¹. The rats breathed spontaneously, and the concentration of sevoflurane and oxygen were monitored continuously employing a calibrated Datex side stream analyzer that sampled from inside of the chamber. The previous study (Zhang et al. 2015) demonstrated that 3% sevoflurane exposure for 2 h daily in 3 successive days was safe for maintaining breath spontaneously and physiological parameters in physiological ranges in rat pups due to its characteristics of small respiratory tract irritation and stable hemodynamics. After the daily 2-h sevoflurane exposure, the rat pups were placed in a thermostated chamber to recover and were returned to the mothers when gaining the righting reflex. Rat pups in the CON group were separated from the dams for identical duration of time in same conditions (37 ± 1 °C), (FiO2 50%), except for exposure to sevoflurane. To study the role of GABA₆R in the present study, a subgroup of rats received three injections of the Na⁺–K⁺–2Cl⁻ cotransporter inhibitor, bumetanide (BUM, 1.82 mg/kg, intraperitoneally, i.p.), 15 min prior to the daily sevoflurane exposure (SEV + BUM group). BUM in this concentration/dose range is commonly used as the most selective and effective of currently available inhibitors of NKCC1 (Tyzio et al. 2014; Dzhala et al. 2005). In order to control for the injections of bumetanide prior to anesthesia, other experimental groups received equal volumes of vehicle (VEH, i.p.) saline at P5, P6, and P7. In order to test whether sevoflurane exposure at different developmental brain stage contributed to different results, juvenile rats receiving identical sevoflurane exposures on P25, P26, and P27 were explored.

**Stress Exposure in Fear Conditioning Training and Fear Extinction Training**

The inescapable electric foot shock stress in fear conditioning training based on the previous study was performed in the present study [19]. Briefly, the conditioning chamber consisted of a brighty lit plexiglass box with stainless steel-grid floor (Shanghai Softmaze Information Technology Co. Ltd., Shanghai, China). An infrared video camera mounted on the rear wall of the cabinets recorded the activity of each rat inside the chamber. The electric foot shock stress in fear conditioning training occurred on P14 for neonatal sevoflurane-exposed rats or on P34 for juvenile sevoflurane-exposed rats. Fear conditioning experiment include a two min exploration period followed by ten conditioned stimulus (CS)–unconditioned stimulus (US) pairings (CS, 70 dB white noise, 20 s duration; US, 2.0 mA electric foot shock intensity, 4 s duration; US was delivered during the last 4 s of the CS presentation) separated by 1 min each. Four days after the conditioning trial, the activities in open-field test were recorded. The fear conditioning acquisition was determined by receiving 2-min CS presentations in an alternative context with distinct visual and tactile cues after a 2 min adaptation period, and then a fear extinction training (for the same set of rats) was performed by receiving 6 × 2 min CS presentations (interval of 2 min). The rate of the rat’s freezing response (defined as the absence of movement except that required for respiration) for every 2 min CS presentation was recorded to measure the fear memory. The level of nonspecific freezing provoked by the new context was controlled for 2 min before the presentation of the cue in that new context. Rats were tested for extinction recall by receiving 2 min CS presentation 48 h after extinction training. At the end of testing for each rat, the arena was cleaned with 75% alcohol to avoid the presence of olfactory cues.

**Open-Field Test**

The activities were determined in the open-field test at P18 for neonatal sevoflurane-exposed rats and P38 for juvenile sevoflurane-exposed rats. In brief, the rats were placed in the center of the open-field chamber (100 × 100 × 40 cm). The rat activities were automatically recorded by a video tracking system and were assessed by measuring the total distance travelled during a 10-min period. At the end of testing for each rat, the arena was cleaned with 75% alcohol to avoid the presence of olfactory cues.

**Measurement of Serum Corticosterone**

Serum corticosterone was measured using commercial ELISA kits (Cayman Chemical Company, Ann Arbor, MI). In order to study the effect of repeated sevoflurane exposures on stress response to electric foot shock in the fear conditioning training, serum levels of corticosterone under basal and stress condition were measured in blood samples collected from the P14 (for neonatal sevoflurane-exposed rats) or P34 (for juvenile sevoflurane-exposed rats) rats 60 min before the process of fear conditioning training and 5 min after the last unconditioned stimulus (After-US). Blood sampling was done using the “tail clip” method. Specifically, the distal 0.5 mm of the tail was removed using a sterile scalpel blade, and blood was allowed to drain directly into a microcentrifuge tube.
Western Blot

The PFC was collected 6 h after exposed to electric foot shock stress in fear conditioning. Rats were deeply anesthetized with sodium pentobarbital (50 mg kg⁻¹, i.p.) and perfused with phosphate-buffered saline through the left cardiac ventricle. The brains were removed from the skull, put into liquid nitrogen, and then stored at 80 °C until further use. On the day of analysis, the brain tissue was allowed to equilibrate to a temperature of 4 °C. Western blot analysis was performed as previously described (Zhang et al. 2015). For primary antibodies, we used rabbit anti-NKCC1 (1:1000; Cell Signaling, USA), anti-KCC2 (1:1000; Cell Signaling, USA), and activated Cleaved Caspase-3 (1:500; Cell Signaling, USA). Antibody anti-β-actin (1:10 000; Sigma, St. Louis, MO) was used to detect β-actin (42 kDa). The membranes were incubated with horseradish peroxidase-conjugated goat anti-rabbit IgG (1:2000; sc-2004, Santa Cruz Biotechnology, Inc., Santa Cruz, CA). The Image J software (NIH Image, Bethesda, MD) was used for semi-quantification of the bands. Expression of β-actin, determined with β-actin antibody, was used to control for loading differences in the total protein amount.

Statistical Analysis

Data are presented as mean ± SEM and analyzed by the Statistical Product for Social Sciences (version 17.0, SPSS Inc., Chicago, IL, USA). Normal distribution of data was analyzed using the Kolmogorov–Smirnov test. Differences among multiple means were assessed by one-way analysis of variance (ANOVA) or repeated-measures two-way ANOVA followed by post hoc Bonferroni test. Independent Student’s t test was used for single comparisons between two groups. A p value less than 0.05 was regarded as a statistically significant difference.

Results

Repeated Sevoflurane Exposures in Neonatal Rats Increased the NKCC1/KCC2 Ratio in the PFC at P14, Which Was Alleviated by Pretreated with the NKCC1 Inhibitor Bumetanide

The authors investigated whether repeated sevoflurane exposures in neonatal rats altered the expression the NKCC1/KCC2 ratio in the PFC during the day of fear conditioning training at P14. The results showed repeated sevoflurane exposures in neonatal rats at P5–P7 which had no statistically difference on expression of NKCC1 (F(2, 15) = 1.21, p = 0.11; one-way ANOVA; Fig. 2A) and KCC2 levels in the PFC at P14 (F(2, 15) = 0.94, p = 0.08; one-way ANOVA; Fig. 2B). However, the resulting NKCC1/KCC2 ratio in the PFC at P14 was significantly increased for neonatal sevoflurane-exposed rats (F(2, 15) = 3.27, p < 0.05; Fig. 2C), which was alleviated by pretreated with the NKCC1 inhibitor bumetanide (F(2, 15) = 3.27, p < 0.05; Fig. 2C). These results suggested the possibility the brain depolarizing GABAAR activity might be altered when exposed to the electric foot shock stress in fear conditioning training at P14 for neonatal sevoflurane-exposed rats, which could be attenuated by pretreated with the NKCC1 inhibitor bumetanide.

Repeated Exposures to Sevoflurane in Neonatal Rather Than Juvenile Rats Increased the Stress Response to Electric Foot Shock in the Fear Conditioning Training

In order to assess whether neonatal exposure to sevoflurane alters the susceptibility to stress exposure, serum corticosterone levels under basal and stress conditions were measured in blood samples collected from P14 rats 60 min before the process of fear conditioning and 5 min after the
last unconditioned stimulus (After-US). The results showed neonatal sevoflurane-exposed had unaltered secretion of corticosterone under the basal condition (Fig. 3A); however, the levels of corticosterone under stress condition for neonatal sevoflurane-exposed rats were significantly higher than CON + VEH group ($F_{(3,21)} = 18.17, p < 0.05$, one-way ANOVA; Fig. 3A). Pretreatment of neonatal rats prior to each sevoflurane exposure with BUM significantly attenuated the heightened secretion of corticosterone after the stress exposure in fear conditioning training ($F_{(3,21)} = 18.17, p < 0.05$; Fig. 3A).

To test whether sevoflurane exposure at different developmental brain stage affect the stress response, juvenile rats receiving identical sevoflurane exposures on P25, P26, and P27 were explored. The results showed there were no statistical difference between SEV + VEH and CON + VEH groups in secretion of corticosterone under basal and stress conditions (Fig. 3B) at P34.

These results indicated repeated exposures to sevoflurane in neonatal rather than juvenile rats increased the neuroendocrine response to future stress exposure, which might be associated with the neonatal enhanced brain depolarizing GABA$_{A}$R activity.

**Repeated Sevoflurane Exposures in Neonatal Rats Increased the Brain Vulnerability to Future Stress Exposure**

The authors tested whether repeated sevoflurane exposures in neonatal rats increased the brain vulnerability to the future stress exposure and whether the adverse results could be attenuated by bumetanide. The basal neuroapoptosis for neonatal sevoflurane-exposed rats were determined without receiving electric foot shock stress at P14. Other independent rat groups were used in determination of stress-induced neuroapoptosis in the PFC 6 h after exposed to electric foot shock stress in fear conditioning training at P14, which was alleviated by pretreated with the NKCC1 inhibitor bumetanide (C). Repeated sevoflurane exposures for juvenile rats at P25–27 had no significant effects on neuroapoptosis in the PFC 6 h after exposed to stress in fear conditioning training at P34 (D). (mean ± SEM; $n=6$ rats/group; *$p \leq 0.05$ vs. CON + VEH group; #*$p \leq 0.05$ vs. SEV + VEH group, one-way ANOVA)
to electric foot shock stress in fear conditioning training at P14. The results showed there was no significantly difference in the neuroapoptosis in the PFC for rats not receiving electric foot shock stress ($p = 0.15$; Fig. 4). However, repeated sevoflurane exposures in neonatal rats at P5, 6, 7 increased the neuroapoptosis in the PFC 6 h after exposed to electric foot shock stress in fear conditioning training at P14 ($F_{(2,15)} = 5.12$, $p < 0.01$; Fig. 3C), which was alleviated by pretreated with the NKCC1 inhibitor bumetanide ($F_{(2,15)} = 5.12$, $p < 0.05$; Fig. 3C). Repeated sevoflurane exposures for juvenile rats at P25–P27 had no significant effects on neuroapoptosis in the PFC 6 h after exposed to stress in fear conditioning training at P34 (Fig. 3D).

These results indicated repeated exposures to sevoflurane in neonatal rather than juvenile rats increased the brain vulnerability to adverse post-stressful factors, which might be associated with the neonatal enhanced brain depolarizing GABA_A activity.

**Neonatal Rather Than Juvenile Sevoflurane-Exposed Rats Exhibited Deficits in Fear Extinction Training and Recall**

For neonatal sevoflurane-exposed rats, there was no significantly difference in the total distance travelled in open-field test among CON + VEH, SEV + VEH, and SEV + BUM groups at P18 (Fig. 5A). The fear conditioning acquisition was determined after the open-field test. One-way ANOVA analysis revealed there were no group differences in pre-CS freezing prior to CS-presentation ($F_{(2,33)} = 0.63$, $p = 0.79$; Fig. 5C). The results showed the rate of freezing under CS-presentation at P18 was significantly increased for CON + VEH, SEV + VEH, and SEV + BUM groups (Fig. 5C). Although the SEV + VEH group exhibited higher rate of freezing than CON + VEH and SEV + BUM in fear conditioning acquisition, there was no statistical difference among CON + VEH, SEV + VEH, and SEV + BUM groups ($F_{(2,33)} = 0.86$, $p = 0.13$; Fig. 5C; one-way ANOVA). For juvenile sevoflurane-exposed rats, there was no significantly difference in the total distance travelled in open-field test between CON and SEV groups at P38 (Fig. 5B). The fear conditioning acquisition at P38 was significantly increased for CON and SEV groups at P38 (Fig. 5D).

Following the determination of fear conditioning acquisition, the authors performed the fear extinction training for neonatal sevoflurane-exposed rats, which was the laboratory basis of exposure therapy for anxiety disorders. Regarding the rate of freezing change during extinction training, there was a significant effect of extinction trial ($F_{(5,165)} = 24.43$, $p < 0.01$; Fig. 6A). Repeated measures two-way ANOVA showed there was significant effect of treatment ($F_{(2,165)} = 18.87$, $p < 0.01$; Fig. 6A). The post hoc Bonferroni test showed SEV + VEH rats had a higher rate of freezing response than the CON + VEH rats at CS3, CS4, CS5, and CS6. Pretreated with the NKCC1 inhibitor bumetanide before neonatal sevoflurane exposures lowered the rate of freezing response in fear extinction training compared with SEV + VEH rats at CS5 and CS6.

The CON + VEH and SEV + BUM rats exhibited comparable, low CS-elicited freezing at CS5 and CS6 in fear extinction training, which suggested the success in retention of fear extinction for both groups (Fig. 6A); however, SEV + VEH rats still exhibited higher CS-elicited freezing compared with the CON + VEH and SEV + BUM groups at CS5 and CS6, suggesting the deficit in retention of fear extinction. Then rats were tested for extinction recall at P20 by receiving 2-min CS presentation. All groups exhibited comparable, low, pre-CS freezing during the adaptation period prior to extinction recall ($F_{(2,33)} = 1.28$, $p = 0.78$; Fig. 6B); however, the groups differed significantly in CS-elicited freezing ($F_{(2,33)} = 16.21$, $p < 0.01$; Fig. 6B). The post hoc Bonferroni test showed that the SEV + VEH rats exhibited significantly higher rate of freezing response compared with the CON + VEH group ($F_{(2,33)} = 16.21$, $p < 0.01$; Fig. 6B) in extinction recall. Pretreated with the NKCC1 inhibitor bumetanide before neonatal sevoflurane exposures reduced the rate of freezing response in fear extinction recall ($F_{(2,33)} = 16.21$, $p = 0.019$; Fig. 6B).

For juvenile sevoflurane-exposed rats, there was no difference between the two groups in pre-CS freezing prior to extinction training ($p = 0.78$; Fig. 6C). Regarding the rate of freezing change during extinction training, there was significant effect of extinction trial ($F_{(5,110)} = 16.23$, $p < 0.01$;...
However, repeated measures two-way ANOVA showed there was no effect of group (F(1,110) = 0.96, p = 0.67; Fig. 6C) and trial-by-group interaction (F(5,110) = 1.49, p = 0.39; Fig. 6C), indicating that both groups exhibited comparable rates of extinction (Fig. 6C). Both groups exhibited comparable, low, pre-CS freezing during the adaptation period prior to extinction recall (p = 0.57; Fig. 6D). There was also no difference between the two groups in CS-elicited freezing (Fig. 6D).

**Discussion**

The present study tested the hypothesis that repeated sevoflurane exposures in neonatal rats increased the brain vulnerability to future stress exposure and resulted in fear extinction deficit and investigated whether the neonatal brain depolarizing GABA$_A$R was involved in mediating these abnormalities. The results showed that repeated sevoflurane exposures in neonatal rats increased the NKCC1/KCC2 ratio in the PFC at P14. Repeated exposures to sevoflurane in neonatal rather than juvenile rats increased the stress response and exacerbated neuroapoptosis in the PFC after exposed to electric foot shock stress in fear conditioning training.

Neonatal sevoflurane-exposed rats exhibited deficits in the fear extinction training and recall. Pretreatment of neonatal rats prior to sevoflurane exposures with bumetanide reduced the NKCC1/KCC2 ratio during the stress exposure in the fear conditioning training and ameliorated most of the subsequent adverse effects. Our study indicate that repeated sevoflurane exposures in neonatal rats increased the brain vulnerability to future stress exposure and resulted in fear extinction deficit, which might be associated with the neonatal enhanced brain depolarizing GABA$_A$R activity.

During the early brain development, neocortical neurons experience activity-dependent, caspase-3-mediated neuroapoptosis. Programmed cell death is an indispensable process for the normal development of the nervous system, which could be activated by physiological or pathological stimuli. In the immature neurons for the developing brain, neuronal apoptosis could be induced by excessive activation of GABA$_A$R (Ikonomidou et al. 2001). There is a high expression level of caspase-3 in the neocortex during the period of early postnatal life and a subsequent decline to low levels during the adulthood. In the present study, there is no significantly difference in the neuroapoptosis in the PFC for rats not exposed to electric foot stress between CON and SEV groups; however, repeated exposures to sevoflurane in
neonatal rats increased the neuroapoptosis in the PFC 6 h after exposed to electric foot shock stress in fear conditioning training at P14, suggesting the increased susceptibility of brain to stress-induced brain injury for neonatal sevoflurane-exposed rats. Repeated sevoflurane exposures in neonatal rats might contribute to disturbance in brain network activity and interfere with the activity-dependent regulation of developmental apoptosis, which eventually increased the brain vulnerability to future stress exposure and resulted in neurological dysfunction.

Disturbing NKCC1/KCC2 balance would result in brain circuit hyperexcitability and contribute to a series of stress-related neurocognitive and psychosocial dysfunctions (Liu et al. 2020; Ben-Ari 2017; Yang et al. 2018; Ju et al. 2018; Cabrera et al. 2020). The peak expression of NKCC1 in rodents is around P5–P7 (Dzhala et al. 2005), which provide the driving force for GABA<sub>A</sub>R-mediated depolarizing response in immature neurons (Dzhala et al. 2005; Ben-Ari 2017; Pfefef et al. 2009). NKCC1 was reported to play a critical role in regulating the function of brain GABAergic transmission and modulating the maternal separation-induced susceptibility to future stress (Hu et al. 2017). Since GABA<sub>A</sub>R-mediated suppression of the activity of limbic-hypothalamic–pituitary–adrenal (LHPA) axis was an important mechanism of adaption to adverse psychosocial stress for the mature brain (Mody and Maguire 2012; Crowley and Girdler 2014), the GABA<sub>A</sub>R-based suppression of LHPA to psychosocial stress in the mature brain might exhibit an opposite, stress-exacerbating effect in the neonatal immature brain. In the present study, the NKCC1/KCC2 ratio was increased in the PFC for neonatal sevoflurane-exposed rats P14, and the neuroendocrine response and brain vulnerability to stress exposure in the fear conditioning training were significantly increased for neonatal sevoflurane-exposed rats, which was alleviated by administration of NKCC1 inhibitor bumetanide before neonatal sevoflurane exposures. We proposed the mechanism that the enhanced depolarizing GABA<sub>A</sub>R activity after neonatal repeated sevoflurane exposures at P5–P7 contributed to the dysregulation of the neuroendocrine response and increased the brain vulnerability to stress exposure later in life. The alleviating effects of bumetanide for neonatal sevoflurane exposed rats are that bumetanide facilitates adaptation of the neonatal brain to adverse stressful stimuli by reversing GABA-initiated signaling from depolarization to inhibition.

Fear extinction provides the experimental basis for fear inhibition, which is the common psychological therapy of anxiety disorders. The anxiety in fear extinction is usually characterized by lacking in the ability to inhibit fear. Fear extinction involves reciprocal PFC interactions with thalamic, hippocampal, and neocortical pathways (Quirk et al. 2006). Recall of fear extinction, which is regarded as...
effective aid in recovery from a tremendous psychologically traumatic event, is demonstrated to be deficient in PTSD. In the current study, repeated exposures to sevoflurane in neonatal rats increased the brain vulnerability to future stress exposure and resulted in fear extinction deficit, which might be associated with paradoxical hyperexcitatory GABA_{A}R activity during neonatal life.

Conclusion

In the present study, our findings indicated that repeated sevoflurane exposures in neonatal rats increased the brain vulnerability to future stress exposure and resulted in fear extinction deficit, which might be associated with the neonatal enhanced brain depolarizing GABA_{A}R activity. The present findings alert the adverse stressful factors after subject to early life sevoflurane exposure and suggest a new potential target in prevention of neonatal anesthesia-related neurocognitive dysfunctions.

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Author Contribution Study design: Bin Liu, Ben-zhen Chen. Experimentation: Wen-qin Zhou, Yu-chao Shang, Fang Li. Data analysis: Li-hua Jiang, Ling Tan. Overall project supervision: Bin Liu. Final manuscript preparation: Ben-zhen Chen, Li-hua Jiang. All authors reviewed the manuscript.

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Availability of Data and Material The data that support the findings of this study are available from the corresponding author, Bin Liu, upon reasonable request.

Declarations

Ethics Approval The present study was approved by the Ethics Committee of West China Hospital, Sichuan University, and was conducted in accordance with the ARRIVE guidelines and the Guide for the Care and Use of Laboratory Animals from the National Institutes of Health (Bethesda, MD, USA).

Consent to Participate Not applicable.

Consent for Publication All authors have seen and approved the manuscript to be published and contributed intellectually to this work.

Conflict of Interest The authors declare no competing interests.

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