Phenotypic dysregulation of microglial activation in young offspring rats with maternal sleep deprivation-induced cognitive impairment

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Despite the potential adverse effects of maternal sleep deprivation (MSD) on physiological and behavioral aspects of offspring, the mechanisms remain poorly understood. The present study was intended to investigate the roles of microglia on neurodevelopment and cognition in young offspring rats with prenatal sleep deprivation. Pregnant Wistar rats received 72 h sleep deprivation in the last trimester of gestation, and their prepuberty male offspring were given the intraperitoneal injection with or without minocycline. The results showed the number of Iba1 microglia increased, that of hippocampal neurogenesis decreased, and the hippocampus-dependent spatial learning and memory were impaired in MSD offspring. The classical microglial activation markers (M1 phenotype) IL-1β, IL-6, TNF-α, CD68 and iNOS were increased, while the alternative microglial activation markers (M2 phenotype) Arg1, Ym1, IL-4, IL-10 and CD206 were reduced in hippocampus of MSD offspring. After minocycline administration, the MSD offspring showed improvement in MWM behaviors and increase in BrdU/DCX cells. Minocycline reduced Iba1 cells, suppressed the production of pro-inflammatory molecules, and reversed the reduction of M2 microglial markers in the MSD prepuberty offspring. These results indicate that dysregulation in microglial pro- and anti-inflammatory activation is involved in MSD-induced inhibition of neurogenesis and impairment of spatial learning and memory.

The stress exposure in the prenatal period can eventually lead to multiple behavioral abnormalities in the adult offspring⁸. Sleep deprivation is a form of physical and emotional stressors during pregnancy, which is associated with several harmful consequences to the mothers and their children, and can damage the mother-infant relationship⁹. In rodent animals, maternal sleep deprivation (MSD) reduces adrenal weight and susceptibility to harmful agents⁵ and causes hypertension and renal abnormalities in pups⁶. Sleep restriction during pregnancy may cause behavioral and oxidative metabolic changes in male offspring⁷. The investigation revealed that more than two-thirds of pregnant women reported poor sleep quality and shortened sleep duration, especially in the third trimester⁸. However, little research has been done to directly examine how MSD alters the physiological and behavioral processes in offspring, and what underlies the alterations.

When a mother is deprived of sleep, widespread alterations in behavioral, cognitive, immune, and metabolic processes can be observed⁹, independent of hypothalamic-pituitary-adrenal (HPA) axis activation⁸. Maternal stress has a profound influence on fetal development and exerts long-time effects on health outcomes in offspring⁸. Recent research showed that pro-inflammatory cytokines, including interleukin 1 (IL-1) and IL-1 receptor, and markers of activated microglia, were elevated in the hippocampus and cerebral cortex of offspring after pregnant stress, suggesting an activation of the local inflammatory response⁹. Prenatal stress also alters microglial development and distribution in postnatal rat brain¹⁰. Fetal brain development is sensitive to changes in glial cell function¹¹, in which permanently activated or “primed” microglia contribute to the long-lasting impairment of adult neurogenesis¹².

Microglia constitute the first line of defense against pathological changes within the central nervous system (CNS) microenvironment¹³ and play important roles in various brain pathologies including psychiatric disor-
Microglial activation is often divided into two phenotypic processes, such as proliferation, differentiation, migration, and survival. Microglial activation is often divided into two phenotypic processes: the classical activation (M1) and the alternative activation/deactivation state (M2). Classically activated microglia may contribute to the reduction of neurogenesis and dysfunction of the neurotrophic system by releasing inflammatory mediators, including cytokines such as Tumor Necrosis Factor-α (TNF-α), interferon-γ (IFN-γ), IL-1β, and IL-6. The alternative phenotype, sometimes called neuroprotective microglial phenotype, is important for antagonizing the inflammatory-induced damages in CNS. Once over-activated in embryonic period, microglia continue to be primed and influence neuron survival into adulthood.

Minocycline is a derivative of tetracycline that exerts neuroprotective properties in neurodegenerative diseases. It is a highly lipophilic molecule capable of crossing the blood-brain barrier. The neuroprotective effect of minocycline has been shown to be mediated by inhibition of the proliferation and activation of microglia. Minocycline administration improved MSD-induced changes in hippocampus-dependent spatial learning and memory of young offspring. Minocycline administration improved MSD-induced changes in hippocampus-dependent spatial learning and memory of young offspring. Hippocampus-dependent spatial learning and memory was assessed using the MWM in young offspring on postnatal Day 21. The four groups did not differ significantly in latency to locate the platform on Day 1. The MSD-saline animals spent more time reaching the hidden platform than the control-saline and control-minocycline groups on Day 2–4. Nevertheless, after minocycline administration, the MSD group had a shorter latency than the MSD-saline group on Day 3–4 (Fig. 1a; day 1: F(3, 26) = 1.844, p = 0.147; day 2: F(3, 26) = 3.458, p = 0.021; day 3: F(3, 26) = 4.600, p = 0.006; day 4: F(3, 26) = 3.039, p = 0.038). When the platform was removed for the probe trial on Day 5, the MSD-saline group spent significantly less time in the training quadrant than other groups (Fig. 1b; F(3, 26) = 3.54, p = 0.018). To further characterize memory deficits, we also recorded the number of platform crossings in a circular area circumscribing the original platform location, and found that the number of platform crossings was significantly smaller in the MSD-saline group than in control groups (Fig. 1c; F(3, 26) = 6.45, p = 0.008). The MSD group after minocycline administration spent more time in the training quadrant than the MSD-saline group. The number of platform crossings in MSD-minocycline offspring was larger than in MSD-saline group. In the reverse trial, when the platform was placed in the quadrant opposite to the original, the latency to locate the platform increased in the MSD-saline group on Day 6 and Day 7.

Results
Minocycline administration improved MSD-induced changes in hippocampus-dependent spatial learning and memory of young offspring. Hippocampus-dependent spatial learning and memory was assessed using the MWM in young offspring on postnatal Day 21. The four groups did not differ significantly in latency to locate the platform on Day 1. The MSD-saline animals spent more time reaching the hidden platform than the control-saline and control-minocycline groups on Day 2–4. Nevertheless, after minocycline administration, the MSD group had a shorter latency than the MSD-saline group on Day 3–4 (Fig. 1a; day 1: F(3, 26) = 1.844, p = 0.147; day 2: F(3, 26) = 3.458, p = 0.021; day 3: F(3, 26) = 4.600, p = 0.006; day 4: F(3, 26) = 3.039, p = 0.038). When the platform was removed for the probe trial on Day 5, the MSD-saline group spent significantly less time in the training quadrant than other groups (Fig. 1b; F(3, 26) = 3.54, p = 0.018). To further characterize memory deficits, we also recorded the number of platform crossings in a circular area circumscribing the original platform location, and found that the number of platform crossings was significantly smaller in the MSD-saline group than in control groups (Fig. 1c; F(3, 26) = 6.45, p = 0.008). The MSD group after minocycline administration spent more time in the training quadrant than the MSD-saline group. The number of platform crossings in MSD-minocycline offspring was larger than in MSD-saline group. In the reverse trial, when the platform was placed in the quadrant opposite to the original, the latency to locate the platform increased in the MSD-saline group on Day 6 and Day 7.
Minocycline administration ameliorated MSD induced deficits in neurogenesis in hippocampus of young offspring. 

BrdU and DCX cells indicated in the DG of hippocampus (Fig. 2a). The total number of BrdU+ cells (b), the number of BrdU+/DCX+ cells (c) and the percentage of BrdU+/DCX+ out of all BrdU+ cells (d) were reduced in the MSD young offspring rats. After minocycline treatment, the total number of BrdU+ cells, the number of BrdU+/DCX+ cells and the percentage of BrdU+/DCX+ out of all BrdU+ cells were increased. Representative figures of the volumes of DG (e) and the GCL areas (f). The volume of DG was not altered in young offspring rats. (h) MSD-mino did not change the volume of GCL subregion in young offspring. * P < 0.05, ** P < 0.01 vs. the control-saline. * P < 0.05, ** P < 0.01 vs. the MSD-saline. Values are the mean ± SEM. Scale bars: A: 20 μm; E, F: 10 μm.

Minocycline administration ameliorated MSD induced deficits in neurogenesis in hippocampus of young offspring. BrdU+ and DCX+ cells indicated in the DG of hippocampus (Fig. 2a). The total number of BrdU-positive cells in the DG of hippocampus in the MSD-saline group was significantly smaller than that of the other three groups. After minocycline administration, however, the number of BrdU-positive cells increased in the MSD group (Fig. 2b; F (3, 18) = 18.49, p = 0.001). The number of BrdU+/DCX+ cells decreased in the MSD group, but increased after minocycline injections (Fig. 2c; F (3, 18) = 135.50, p < 0.001). While the ratio of BrdU+/DCX+/BrdU+ decreased in the MSD-saline group, it almost reached the control level after minocycline administration (Fig. 2d; F (3, 18) = 46.18, p < 0.001).

DAPI stained sections were used to measure the volumes of neurogenesis regions in the offspring hippocampus. Fig. 2e outlines DG and Fig. 2f shows the subregion of GCL. No significant differences in the volumes of DG (Fig. 2g) and GCL areas (Fig. 2h) were identified between the MSD-saline and MSD-minocycline groups (Fig. 2g; DG: F (3, 18) = 6.138, p = 0.062; Fig. 2h; GCL: F (3, 18) = 4.565, p = 0.720).

Minocycline treatment reduced microglial activation caused by MSD in hippocampus of young offspring rats. The expression of mRNA for Iba1 increased in the MSD-saline young offspring. After minocycline administration, the level of Iba1 was reduced (Fig. 3a; F (3, 26) = 7.542, p = 0.002). The control-saline offspring exhibited no
activation of microglia. The microglia in the MSD-saline offspring had large somas, short thick processes, and rounded amoeboid morphologies, known as hallmarks of activated microglial cells. The morphology of microglia in the MSD-minocycline was rod-shaped cell bodies with fine, ramified processes, a resting condition (Fig. 3b). Compared with MSD-saline group, the number of Iba1 positive microglia in the hippocampus of the MSD-minocycline group was significantly reduced (Fig. 3c; \( F(3, 18) = 531.232, p = 0.005 \)).

Minocycline decreased the expression of M1 markers in the hippocampus of young offspring rats. The expression of M1 markers IL-1β, IL-6, TNF-α, CD68, and iNOS was increased in hippocampus of the MSD offspring. These increases were significantly attenuated by treatment with minocycline (Fig. 4a: \( F(3, 26) = 4.573, p = 0.011 \); Fig. 4b: \( F(3, 26) = 8.469, p = 0.001 \); Fig. 4c: \( F(3, 26) = 9.012, p = 0.001 \); Fig. 4d: \( F(3, 26) = 7.571, p = 0.015 \); Fig. 4e: \( F(3, 26) = 6.202, p = 0.016 \)). By contrast, the expression of M2 markers (Arg1, Ym1, IL-4, IL-10, and CD206) was reduced in the hippocampus of the MSD young offspring. After minocycline administration, the expression of Arg1, IL-4, IL-10, and CD206 was enhanced in the MSD offspring (Fig. 5a: \( F(3, 26) = 6.965, p = 0.002 \); Fig. 5b: \( F(3, 26) = 11.867, p = 0.012 \); Fig. 5c: \( F(3, 26) = 11.219, p = 0.001 \); Fig. 5d: \( F(3, 26) = 8.391, p < 0.001 \)). However, Ym1 mRNA level was not changed by minocycline treatment in the MSD offspring (Fig. 5b; \( F(3, 26) = 18.907, p < 0.001 \)).

Minocycline attenuated IL-6 and increased Arg1 protein expression derived from activated microglia in the hippocampus of young offspring rats. The expression of M1 marker IL-6 was higher and that of M2 marker Arg1 was lower after MSD in the hippocampus of prepuberty offspring. IL-6 mRNA level increased 37 fold in the MSD-saline offspring over the control-saline offspring, whereas Arg1 mRNA level was reduced by 90% (Figs. 4b and 5a). The IL-6 and Arg1 protein were detected using ELISA kits. In the hippocampus, the expression of IL-6 was enhanced after MSD compared with the control. The increase was attenuated by
minocycline treatment (Fig. 6a; $F_{(3, 26)} = 9.780, p = 0.006$). The quantity of Arg1 protein increased in the MSD-minocycline group compared to the MSD-saline group (Fig. 6b; $F_{(3, 26)} = 8.751, p = 0.007$). IL-6 and Arg1 were generated from microglia of the MSD-minocycline group in the hippocampus of offspring rats (Fig. 6c, 6d).

**Discussion**

Our data provide evidence that microglia play a pivotal role in modulating the impact of MSD on hippocampus neuronal activity and hippocampus-regulated behavior in the offspring. Minocycline can dampen MSD-induced microglial activation and hippocampal neu-
rogenesis impairment, and also reverse the spatial learning and memory deficits in the MSD young offspring. The neuroprotective effect of minocycline in MSD offspring rats was related to switching the pro-inflammatory microglial response to anti-inflammatory microglial phenotype.

Stress during pregnancy has lasting effects on neurodevelopment and increases vulnerability to psychopathology in offspring. In this study, we investigated the effects of sleep deprivation during pregnancy on the neurogenesis of the prepuberty male offspring. The total numbers of BrdU/cells and BrdU+/DCX+ cells in MSD offspring were smaller than that of the control. This showed that hippocampal neurogenesis is reduced in the DG in young rats by MSD, which conforms with the findings of our previous research. These results support the concept that prenatal stress can induce lasting and profound changes in the offspring, potently inhibiting neurogenesis. The low level of neurogenesis was likely due to the established stress hyper-responsiveness, including the inflammatory response or neurodegeneration. We observed the suppressed neurogenesis in MSD offspring restored by intraperitoneal (i.p.) administration of minocycline, which was associated with specific inhibition of microglial activation. However, no significant differences were identified in the volumes of hippocampus neurogenesis regions in the MSD-minocycline offspring. In addition to new neurons, other factors such as decreases in the survival of neurons, alterations in the somatodendritic, axonal, and synaptic components and glial changes are also corrected with hippocampal volume loss.

Our behavioral studies with the young offspring rats that were prenatally stressed using the MWM tasks demonstrated impairment of spatial working and memory, which requires a fully functional hippocampus. As shown in Fig. 1, the MSD-minocycline offspring spent significantly less time than the MSD-saline groups locating the hidden platform during the task acquisition and the reverse trial, spent more time in the target quadrant and they had a larger number of platform crossings than the MSD-saline offspring in the spatial exploration test. This indicates a deficit reversed in hippocampus-dependent spatial learning and memory in the MSD-minocycline offspring rats. In rodents, the hippocampus has long been recognized as a critical structure for encoding spatial information. MSD-induced impairment of hippocampus-dependent behavioral tasks was connected to inhibition of hippocampal neurogenesis in the offspring rats. According to hippocampal neurogenesis, the spatial learning and memory of MSD offspring were improved after anti-inflammatory treatment with minocycline. It appeared that the anti-inflammatory property of minocycline was responsible for the behavioral effects.

Although Iba1 is constitutively expressed by microglia, it is only moderately expressed in quiescent microglia, but strongly expressed in response to activating stimuli. MSD significantly increased the expression of Iba1 on mRNA level and the density of Iba1 positive cells in the offspring rats, as indicated by immunohistochemistry. We have shown in our previous research that the morphological features...
of microglia were large somas, short thick processes, and a rounded amoeboid in the offspring of MSD. This suggests that sleep deprivation in pregnant dams could induce microglial activation in the prepuberty offspring, similar to prenatal immune challenges. The pro-inflammatory status is considered a risk factor for development of behavioral alterations and cognitive deficits in animals and humans. Minocycline can reduce microglia activation, which was demonstrated in this experiment by the down-regulation of mRNA expression of Iba1 and the reduction in the number of immunolabeling Iba1⁺ cells. The results indicated that the improvements of MWM behaviors and neurogenesis by minocycline treatment in the young offspring rats were related to the inhibition of activated microglia.

Insufficient sleep and poor sleep quality have pro-inflammatory effects, which are independent of nonspecific consequences of sleep deprivation procedures. In this study, the multiple-platform method was used to avoid the stresses of activity restriction. The pro-inflammatory effect of sleep disturbance during pregnancy is a risk factor for embryonic developments through maternal-placental-fetal inflammatory pathways. Intrauterine inflammatory exposure may induce microglial changes and mediate programming of neuroinflammatory processes in offspring. Although cytokines can be produced by astrocytes, endothelial cells, even neuron, microglia are the primary source for inflammatory mediators. The expression of pro-inflammatory factors such as IL-1β, IL-6, TNF-α, and iNOS was increased in the hippocampus of young rats whose mothers experienced sleep deprivation. Meanwhile, the expression of anti-inflammatory cytokines IL-4 and IL-10 was reduced in the MSD prepuberty rats. Opposing the deleterious effects of pro-inflammatory cytokines, anti-inflammatory cytokines such as TGF-β and IL-10 have pro-neurogenic effects on adult neural stem/progenitor cells. The dysregulation between pro- and anti-inflammatory cytokines plays an important role in the impairments of neurogenesis.

Studies on peripheral macrophages have led to the identification of distinct activation profiles, referred to as classical (M1) and alternative (M2) activation states. These are characterized by different markers being expressed and different functions of the cells. On one hand, the expression of M1 markers (IL-1β, IL-6, TNF-α, CD68 and iNOS) gradually increased in the MSD group but not in the control groups. On the other hand, the expression of M2 markers (Arg1, Ym1, IL-4, IL-10, and CD206) was reduced in the MSD-saline group. Our data suggested that the expression of pro-inflammatory cytokines was induced and resulted in M1 microglial activation in the young offspring rats after MSD. Stress induced HPA axis hyperactivity can affect microglial activation and phenotypes, followed by neuroinflammatory outcomes. These results support the concept that prenatal stress can induce long-term pro-inflammatory effects and classical microglial activation or M1 phenotype. With regard to microglial phenotypes, we paid special attention to the current study to the alternative activation of microglia in the young offspring whose mothers were exposed to sleep deprivation. In contrast to the enhanced M1 microglial markers, the expression of M2 markers including Arg1, Ym1, IL-4, IL-10, and CD206 (Fig. 5) were reduced in the MSD offspring rats. Numerous studies have concluded that M2 microglia was neuroprotective while M1 microglia were neurotoxic. M1 microglia may contribute to the deficit of hippocampal neurogenesis and related behaviors by releasing inflammatory mediators, including cytokines such as TNF-α, IL-1β and IL-6, which will ultimately cause local inflammation and neurodegeneration. With minocycline treatment, the M2 markers in MSD offspring were restored to the level of the control rats. The effects of minocycline on microglial activated phenotypes were consistent with its pro-neurogenic effect and neurobehavorial actions in young offspring rats. Results indicated that MSD caused deficits of hippocampus-dependent behavior and reduction of hippocampal neurogenesis in the offspring rats, which strongly correlated with microglial activation phenotype in young offspring rats.

Microglia, the brain’s resident innate immune cells, are thought to derive from myeloid precursor cells that infiltrate the central nervous system via local blood vessels during embryonic and early postnatal life. In the rat, microglia are first seen between embryonic Days 12–14, and then a rapid increase in the number of microglia during late trimester of fetal development period was observed. Microglia are critical for early brain development and can respond vigorously to stress. The third trimester appears to be a particularly sensitive period to inflammatory challenge. Once prenatally challenged, the long-lived microglia are maintained in an activated or primed state into adulthood. Our previous results demonstrated that microglial activation induced by late-stage pregnancy stress could last until postnatal day 21. Microglia constitute the major niche and play a key role in controlling multiple steps of neurogenesis. Prenatal stress induces microglia into a preponderant M1 cytotoxic phenotype in offspring rats. Dysregulated microglial activation leads to the imbalance of pro- and anti-inflammatory cytokines. The pro-inflammatory cytokines in CNS can damage neural cells directly or indirectly. The pro-inflammatory cytokines IL-1β, IL-6, TNF-α, and iNOS were increased and anti-inflammatory cytokine IL-4 and IL-10 was reduced after MSD (Fig. 4). It has been shown that the activation of microglia is associated with a reduction of new neurons in the rodent hippocampus, mainly due to the decreased survival of the new neurons. Minocycline suppressed the production of pro-inflammatory molecules in the MSD prepuberty offspring, improved hippocampus-dependent spatial learning and memory, and promoted hippocampal neurogenesis. Minocycline exerts its neuroprotective effect by inhibiting the microglial polarization into M1, facilitating M2 activation.

Minocycline is a semisynthetic tetracycline that exerts anti-inflammatory effects completely separate from its antimicrobial actions. Studies suggest that it provides neuroprotection because it can cross the blood-brain barrier and inhibit the proliferation and activation of microglia. Minocycline has neuroprotective qualities in animal models of neurodegenerative diseases and has been evaluated in clinical trials for a number of neurodegenerative or autoimmune diseases which affect the CNS, including stroke, Parkinson’s disease, Huntington’s disease, and multiple sclerosis. These neuroprotective activities of minocycline have been attributed to its inhibitory effects on microglial activation. This study indicated that microglial M1 suppressors such as minocycline could have potential benefit for behaviors and neurogenesis of offspring whose mothers experienced sleep deprivation during pregnancy.

In this study, we report for the first time that anti-inflammatory treatment can counteract neuroinflammatory responses and ameliorate cognitive impairment caused by MSD in offspring. Our study indicates that microglia play a significant role in determining neuronal proliferation and survival, which are responsible for the impairment of spatial learning and memory of young offspring rats in MSD model. The M1 microglial activation profiles with prominent pro-inflammatory cytokines are relevant to neural defense and neuropathology. Minocycline inhibit M1-biased microglial response and have beneficial effects in the young offspring rats whose mothers experience sleep deprivation during pregnancy.

Methods
All experimental procedures were approved by the Institutional Animal Care and Use Committee at the University of Electronic Science and Technology of China. The methods were carried out in accordance with the approved guidelines.

Animals. Adult male and female Wistar rats (three months old) were purchased from Chengdu Dossy Biological Technology Co., Ltd. (Chengdu, China), kept individually under standard housing conditions (12/12 h light/dark phase, lights on at 7:00 A.M., humidity 55 ± 5%, temperature 23–25 °C), with ad libitum access to food and water, and habituated to the animal facilities for one week. The female rats were subjected to a timed mating procedure as described previously. The first day of pregnancy was defined as gestational day (GD) 1.
Sleep Deprivation. Sleep deprivation was performed using modified multiple-platform methods on late trimester (GD 18). Pregnant Wistar rats (n = 25) were placed individually in the experimental device in previously described26. All platform exposure began at 10:00 a.m. and ended after 72 h when the pregnant rats were put back into their home cages. All rats had free access to water and food throughout the experiment.

Minocycline treatment. After weaning (21 days old), 2–3 male offspring were selected from each mother group. The male prepuberty offspring from non-MSD and MSD were given l.p. injections of saline and minocycline (50 mg/kg, Sigma-Aldrich, St. Louis, MO, USA) twice daily (8:00 a.m. and 8:00 p.m.) for the first two days, and then i.p. injections once daily at 8:00 a.m. for the following three days. Minocycline dose was selected according to the established dose regimen27. Minocycline was diluted in saline. On the second day after five days of injections, Morris Water Maze, Immunohistochemistry, ELISA and Real-Time PCR were performed among offspring. To avoid the effects of hormonal variation in females, only male offspring were used for the study. The offspring were divided into four groups: control–saline (n = 27), MSD–saline (n = 27), control–minocycline (n = 26), and MSD–minocycline (n = 28) groups were used in the MWM experiment.

**Morris Water Maze.** The prepuberty offspring rats (21 days old) were to find a hidden platform (9 cm in diameter) 2 cm under the water surface in a pool (1 m in diameter), four trials per day for four consecutive days as previously described28. All MWM testing was performed between 9 a.m. and 2 p.m. The MWM was performed by a researcher blinded to the identity of the groups. The young offspring rats of the control-saline (n = 8), MSD-saline (n = 8), control-minocycline (n = 7), and MSD-minocycline (n = 8) groups were used in the MWM experiment.

**Immunohistochemistry.** Offspring rats (21 days old) from the control-saline (n = 5), MSD-saline (n = 5), control-minocycline (n = 5), and MSD-minocycline (n = 5) groups were measured in the immunohistochemical analysis for neuronal proliferation and microglial morphology. Prepuberty offspring rats were given one injection of Bromodeoxyuridine (BrdU) (50 mg/kg; Sigma, USA) 48 h before the experiment.

Minocycline treatment experiment. Minocycline was stored at 4°C and collected in PBS with 0.03% sodium azide. Six sequential slices were placed into each well of a 12-well plate and stored at 4°C for 24 h. Perfusion was performed transcardially with saline, followed by 4% paraformaldehyde in phosphate-buffered saline (PBS). Tissues were transversely cut into coronal sections (35 μm thickness of brain slice and number of sections54). The volumes were measured using software ImageJ, version 1.45J (National Institutes of Health, Bethesda, MD, USA). Iba1 positive stain and cell morphology. Positive cells were manually counted under a 40× objective fluorescence microscope (Olympus BX51). Photomicrographs were taken with a high-resolution digital camera. 

**Cavalieri principle.** The volumes were measured using software ImageJ, version 1.45J (National Institutes of Health, Bethesda, MD, USA). Iba1 was used to evaluate the microglial activated phenotype according to the molecular mechanisms. To quantify the proliferating cells and microglia, every sixth section from one group was used in the immunohistochemical analysis for neuronal proliferation and microglial morphology. Prepuberty offspring rats were given one injection of Bromodeoxyuridine (BrdU) (50 mg/kg; Sigma, USA) 48 h before the experiment.

**Statistical Analysis.** All data were presented as mean ± SEM. The results were analyzed with repeated measures analysis of variance. Differences were considered significant when p < 0.05. Analyses were conducted using SPSS for Windows® v.17 (SPSS Inc., Chicago, USA).

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Author contributions
Author Z.Y. and Q.Z. designed the study and wrote the protocol. Authors Y.C. and C.P. managed the literature searches and analyses. Authors X.X. and X.W. undertook the statistical analysis. Authors Q.Z., Y.F., J.Z., W.J. and S.Y. did the experiments. Author Q.Z. wrote the draft of the manuscript. All authors reviewed the manuscript.

Additional information
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