Hippocampal neurogenesis plays an important role in the onset and treatment of depressive disorders. Previous studies suggest that paeoniflorin could be used as an antidepressant for treating rats subjected to chronic unpredictable stress. In this study, the effects of paeoniflorin on neurogenesis in the hippocampus dentate gyrus and potential mechanism of action are further investigated in chronic unpredictable stress-induced rat. Results suggest that paeoniflorin markedly increased both sucrose consumption and the number of 5-bromo-2-deoxyuridine-positive cells in the dentate gyrus of chronic unpredictable stress-induced rats, and the ratio of co-expressed 5-bromo-2-deoxyuridine and glial fibrillary acidic protein-positive cells, but exerted no significant effect on the ratio of co-expressed 5-bromo-2-deoxyuridine and neuronal nuclear-positive cells. Compared with the vehicle group, a significant increase was detected in the number of brain-derived neurotrophic factor-positive cells and the expression of brain-derived neurotrophic factor mRNA in the hippocampus of the paeoniflorin-treated group. According to the results, paeoniflorin promoted neural stem cell proliferation, their differentiation into astrocytes, and neurogenesis in the hippocampal dentate gyrus of chronic unpredictable stress-induced rats. Apart from enhancing the protein expression and gene transcription of brain-derived neurotrophic factor, it also activated the expression of tropomyosin receptor kinase B (a high-affinity receptor of brain-derived neurotrophic factor). This suggests that paeoniflorin might promote neurogenesis in the hippocampus dentate gyrus of chronic unpredictable stress-induced rats and act as an antidepressant by regulating the brain-derived neurotrophic factor-tropomyosin receptor kinase B signaling pathway.

Keywords
- Paeoniflorin; antidepressant; chronic unpredictable stress; neurogenesis; dentate gyrus; BDNF-TrkB signaling pathway

1. Introduction
Globally, over 300 million people (equivalent to 4.4% of the world’s population in 2015) are estimated to suffer from depression. Depression is ranked by the World Health Organisation as the single largest contributor to global disability (7.5% of all years lived with disability in 2015) (World Health Organization, 2017). Numerous preclinical and clinical studies have reported that adult neurogenesis in dentate gyrus (DG) is closely related to depression (Drnovsky and Hen, 2006; Cameron and Schoenfeld, 2018). Hippocampal atrophy is observed in patients with depressive disorders and changes in hippocampal volume are possibly caused by lowered DG neurogenesis (Malberg, 2004; Alfonso et al, 2004). Santarelli et al and others (Santarelli et al, 2003; Bulmash et al, 2009) have reported that the behavioral effects of antidepressants are dependent upon hippocampal neurogenesis in animal models of depression.

Hippocampal neurogenesis is regulated by multiple factors in vivo and in vitro, among which brain derived neurotrophic factor (BDNF), with high-affinity binding to tropomyosin receptor kinase B (TrkB), is critical for hippocampal neurogenesis in patients with depressive disorders (Castren and Kojima, 2017; Begni et al, 2017). Its expression reflects the severity of lesions in neurodegenerative diseases (Begni et al, 2017) and its deficiency reduces proliferation of newborn granule cells. Additionally, injected exogenous BDNF increases neurogenesis of granule cells (Hu et al., 2010; Zheng et al, 2012; Wang et al, 2017). Currently, antidepressants are reported to activate the functions of BDNF and TrkB in several neuronal processes, such as excitation, development, apoptosis, and synaptic plasticity (Leal et al, 2017; Li et al, 2018).

Paeoniflorin is a major active ingredient in the root of Paeonia lactiflora Pall. (Ranunculaceae), a Chinese medicinal herb widely used for treating depression-like disorders in many prescriptions of traditional Chinese medicine, including “Danggui Shaoyao San” and “Xiaoyao powder” (Qiu et al, 2013a). In many studies, paeoniflorin is found to be effective in treating several types of animal models of depression. Some animal experiments related to depression have shown paeoniflorin to significantly shorten swimming immobility time and tail suspension of mice, and blepharoptosis induced by reserpine antagonist (Qiu et al, 2013b; Cui, 2009). It also markedly ameliorates depressive-like behaviors in interferon-induced rats; dentate gyrus; BDNF-TrkB signaling pathway; Paeoniflorin; antidepressant; chronic unpredictable stress-induced depressive-like rats with paeoniflorin
α (IFN-α)-induced mice, ovarioctomized rats under stress, and chronic unpredictable stress (CUS) induced rats (Qiu et al., 2013a; Huang et al., 2015; Li et al., 2017). This suggests that paeoniflorin may be useful in the prevention of depression. In in vitro experiments paeoniflorin has been observed to be highly effective in protecting PC12 cells or neurons from damage by some drugs, leading to a significant increase in viable cell number, mortality reduction, and protection of these cells (Mao et al., 2012, 2011, 2010). However, its role in adult neurogenesis have neither been studied nor reported.

This study further investigated the impact of paeoniflorin on neurogenesis in the hippocampus DG of CUS-induced rat and its possible mechanisms of action.

2. Materials and methods

2.1. Reagents and chemicals

Paeoniflorin (purity > 95%, 201110) was obtained from Guizhou Dida Technology Co., Ltd; imipramine (090M14881V), 5-bromo-2-deoxyuridine (BrdU) (HMBC4217TV) and mouse monoclonal BrdU antibody (D32N4389) were obtained from Sigma-Aldrich (Shanghai) Trading Co., Ltd; both rabbit polyclonal glial fibrillary acidic protein (GFAP) antibody (1013292) and rabbit monoclonal neuronal nuclei (NeuN) antibody (1011151) were obtained from Millipore. Tetramethylrhodamine (TRITC)-labelled goat anti-rabbit antibody (101272) and goat anti-mouse antibody (97860), anti-fade mounting medium were obtained from Santa Cruz Biotechnology, Inc. Rabbit anti-rat TrkB polyclonal antibody (E0112) were obtained from Abcam. 2′,7′-dichlorofluorescein diacetate (DCF-DA) (D3999), fluorescein isothiocyanate (FITC)-labelled goat anti-rabbit antibody (101272) and rabbit anti-rat BDNF polyclonal antibody (E0112) were obtained from Sigma-Aldrich (Shanghai) Trading Co., Ltd; 2′,7′-dichlorofluorescein diacetate (DCF-DA) (D3999), fluorescein isothiocyanate (FITC)-labelled goat anti-rabbit antibody (101272) and rabbit anti-rat BDNF polyclonal antibody (E0112) were obtained from Santa Cruz Biotechnology, Inc. Rabbit anti-rat TrkB polyclonal antibody (GR45120-6) was obtained from Abcam. 2× SYBR Premix Ex Taq (B704-1), Taq enzyme (DR100B), M-MLV RTase (M-MLV Reverse Transcriptase, CK3401E), Oligo d(T) 18 Primers (B1801-1), ribonuclease (RNase) inhibitor (CK6301B) and dNTP (Deoxynucleotide, B1801-1) were obtained from Takara Biotechnology (Dalian) Co., Ltd. Gene sequences were obtained from Genbank. Primers were designed using Prime premier 5 and synthesized by Sangon Biotech (Shanghai) Co., Ltd. Chloral hydrate was dissolved in normal saline to give a 10% chloral hydrate solution.

2.2. Experimental animals

Male Sprague Dawley rats (140–160 g, four weeks old) were supplied by the Animal Center of Zhejiang Chinese Medical University (license number SCXK (Zhejiang) 2008-0115). Subjects were housed at room temperature (24 ± 1°C, humidity 50 ± 10%) with a 12:12 hour light/dark cycle (light on 8:00 am) and fed and watered ad libitum for 7 days. Experimental procedures were approved by the Animal Experimentation Ethics Committee of Zhejiang Traditional Medical University (ZSLL-2012-057), and were conducted as specified by the National Institutes of Health Guide for the Care and Use of Laboratory Animals.

2.3. Experimental procedure

A full chronological description of the experimental protocol is given in Fig. 1.

2.4. Preparation of depression model and treatment

Subjects were randomly divided into four groups, including control, vehicle, paeoniflorin-treated (60 mg/kg (Qiu et al., 2013a)) and imipramine-treated (10 mg/kg (Qiu et al., 2013a)), each of which had eight subjects. The CUS animal depression model employed here was modified from a previously reported model (Qiu et al., 2013a). Subjects in stressed groups were exposed for seven weeks to daily varying stressors (9:00 am): 24 hours of food deprivation, 24 hours of water deprivation, 24 hours of cage tilt (45°), one minute of tail pinch (1 cm from tail end), two hours of restraint stress, four minutes of exposure to 45 or 4°C, and overnight illumination. The control group was not stimulated. Three weeks after exposure to stressors, subjects were intraperitoneally injected with paeoniflorin and imipramine at 2 mL/kg (8:30 am) one hour before stress stimuli every day for four weeks. The same volume of normal saline was administered to the control and the vehicle groups.

2.5. Sucrose preference test

The sucrose preference test was performed after seven weeks of CUS (10:30 am) and subjects were adapted by training to sucrose before the test, which was performed as previously described (Qiu et al., 2013a). Briefly, subjects were adapted by training to 1% sucrose solution (w/v): two bottles of solution were placed in each cage, and 24 hours later the solution in one bottle was replaced with tap water for 24 hours. After adaptation, subjects were deprived of water and food for 12 hours and then exposed to the sucrose preference test, where they were free to access either bottle. After two hours, the volumes of sucrose solution and water consumed were recorded and sucrose preference calculated.

2.6. BrdU treatment and sample preparation

Five subjects were randomly taken from each group. On the 4th day of the 7th week of CUS, BrdU (50 mg/kg) was given to the subjects six times via intraperitoneal injection at intervals of 12 hours (Lee et al., 2013). After seven weeks of CUS, subjects received a BrdU injection, were anesthetized, and fixed (4% polyformaldehyde) by perfusion through the left ventricle. Subjects were then immediately decapitated and the brains were fixed overnight (4% polyformaldehyde) for immunofluorescent staining and immunohistochemical tests. Finally, brains were collected 24 hours after dosing so as to test proliferation and differentiation of hippocampus neural stem cells in CUS subjects. Subjects that did not receive a BrdU injection were anesthetized by intraperitoneal injection (10% chloral hydrate), and decapitated. Brains and hippocampi were immediately split on a glass table, weighed, frozen in liquid nitrogen, and used to measure the expression of hippocampus-related mRNA.

2.7. Immunofluorescent staining

BrdU, GFAP, and NeuN expression were measured by immunofluorescent staining, a ratin followed by embedding (Takahashi and Ishiguro, 1991). Coronal sections (5 μm) were continuously taken by paraffin slicing machine (HM 340E Rotary Mi-
crotome, Thermo Scientific Co., Ltd) with one slice chosen from every six slices. After dewaxing (Takahashi and Ishiguro, 1991), antigens were retrieved in 0.01 mol/L sodium citrate buffer (pH = 6.0) and soaked in PBS containing 3% H2O2 to suppress the activity of endogenous peroxidase. Following incubation in 2 mol/L HCl at 37 °C for 50 minutes, BrdU antigens of DNA were unmasked. After placement in 0.1 mol/L NaBH4 at 37 °C for 1 hour. Goat anti-goat FITC secondary antibody (1:200) was added and slices were incubated at 37 °C for 45 minutes. Goat anti-mouse TRITC secondary antibody (1:200) was added, slices were then incubated at 37 °C for 45 minutes and reacted without exposure to light. Anti-rabbit NeuN or GFAP primary antibody (1:500) was added and incubated at 37 °C for one hour. Goat anti-goat FITC secondary antibody (1:200) was added and slices were incubated at 37 °C for 45 minutes. Antifade mounting medium was added and the flask was covered with a glass slide. Observations (559 nm) were conducted by Fluoview TM FV1000 laser scanning confocal microscope (Olympus Co., Ltd). Positive cells in hippocampus DG were counted with Image-Pro Plus.

2.8. Immunohistochemical test

BDNF and TrkB protein expression was measured by immunohistochemical test. Methods for material collection, paraffin embedding, slicing, dewaxing (Takahashi and Ishiguro, 1991), high-pressure antigen retrieval and suppression of endogenous peroxidase activity were the same as for immunofluorescence staining. Either anti-mouse BDNF (1:200) or TrkB (1:200) primary antibody was added and the solution cultured overnight (4°C). Rabbit secondary antibody was incubated for one hour (37°C) and DAB color development then performed (1:200, 2-10 min). Hematoxylin staining (2 min) was followed by cells differentiation (10 s chlorohydric acid). Cells were kept transparent by routine dehydration procedures and mounted in neutral balsams. All immunohistochemical positive cells were counted using Image-Pro Plus.

2.9. qRT-PCR test

The expression of BDNF mRNA was detected by qRT-PCR. The hippocampus was homogenized with liquid nitrogen and Trizol solution added. After addition of chloroform, oscillation, and even mixing, the supernatant layer was taken (0.4 mL). Iso-propanol (0.4 ml) was added and the solution evenly mixed, the mixture was then stood still at room temperature. Sediments were removed after centrifugation and cleaning (500 µL 70% ethanol). The sediment was dissolved (50 µL DEPC-H2O) to obtain the total tissue RNA. Total RNA was converted to cDNA using MMLV reverse transcriptase. For real-time PCR, the SYBR qPCR Real-Time kit was used according to the instructions and the product amplified by Bio-Rad Mini Optico realtime PCR. Amplification conditions were set to a 30-cycle program (94°C, 55°C, 72°C, each for 30 s). GAPDH was used as an internal control for the PCR amplification. Primer sequences for BDNF were: F: 5’-CAGTGGCTGGCTCTCATTAC-3’ and R: 5’-CGAAACAGAAGCACAGAAA-3’. Primer sequences for GAPDH were: F: 5’-CCCACGGCAAGTTCAACGGCA-3’ and R: 5’-TTGCACTTTTCTCCAGGCCG-3’. The Ct value of each sample was determined. The relative expression (2^(-∆∆Ct)) of target genes in each sample was calculated by comparing them with those of the control group and using GAPDH as an internal control.

2.10. Statistical analysis

Data were statistically processed using SPSS (Windows v.13.0) and reported as mean ± standard error of the mean (SEM). One-way analysis of variance (ANOVA) was performed, and intergroup differences were measured by Dunnett’s test, where differences were considered statistically significant for a p-value < 0.05.

3. Results

The effect of paeoniflorin on sucrose consumption by CUS subjects is given in Fig. 2. One-way ANOVA revealed that the percentage of sucrose consumption significantly differed between groups [F(3,31) = 11.5, p = 0.01]. Results suggest the vehicle group exhibited a lower sucrose preference than the control group (p = 0.041) and the paeoniflorin group consumed more sucrose than the vehicle group (p = 0.041). Long-term imipramine treatment
led to recovery of sucrose preference in CUS subjects ($p = 0.005$ vs. CUS–saline group).

The effect of paeoniflorin on BrdU expression in hippocampus DG of CUS-treated subjects is shown in Fig. 3. One-way ANOVA showed BrdU expression to be significantly different between groups [$F(3,19) = 8.7, p = 0.001$]. BrdU-positive cells in the DG appeared to have red nuclei. They were either round or oval-shaped and were detected in isolated cells or clusters (Fig. 2). Compared with the control group, significantly fewer BrdU-positive cells were detected in the hippocampus DG of CUS subjects ($p = 0.012$). The number of BrdU-positive cells in hippocampus DG was significantly increased in paeoniflorin and imipramine treated groups when compared with the vehicle group ($p = 0.002, p = 0.0$, respectively) (Fig. 3E).

The impact of paeoniflorin on the ratio of co-expressed BrdU/GFAP positive cells in the hippocampus DG of CUS subjects is illustrated in Fig. 4. Co-expressed BrdU/GFAP positive cells were mature GFAP-immunoreactive astrocytes that differentiated from BrdU-positive cells. Somas of GFAP-immunoreactive cells (green) were swollen and fat and had many dendrite-like protrusions. The protrusions were thickened and heavily stained. Positive cells were individually or collectively aggregated into clusters. In co-expressed BrdU/GFAP-positive cells, BrdU was expressed inside the nucleus (red), and GFAP was expressed in the cytoplasm (green) and protruded like dendrites. In the control group, GFAP-immunoreactive cells were remarkably extended and long. However, they rarely protruded and were short in the hippocampus of CUS subjects (Fig. 4). One-way ANOVA results suggested that the ratio of co-expressed BrdU/GFAP positive cells significantly differed among groups [$F(3,19) = 3.2, p = 0.054$]. Compared with the control group, the ratio of co-expressed BrdU/GFAP positive cells was significantly smaller in the hippocampus DG of CUS subjects ($p = 0.026$). Compared with the vehicle group, a significant increase was detected in the ratio of co-expressed BrdU/GFAP positive cells in hippocampus DG of paeoniflorin and imipramine treated groups ($p = 0.044, p = 0.025$, respectively), as shown in Fig. 4E.

The effect of paeoniflorin on the ratio of co-expressed BrdU/NeuN positive cells in hippocampus DG of CUS subjects is shown in Fig. 5. Changes to the number of co-expressed BrdU/NeuN positive cells in the hippocampus DG of subjects were measured by double immunofluorescence staining. Co-expressed BrdU/NeuN positive cells were mature NeuN-immunoreactive neural cells into which BrdU-positive cells were differentiated.
The effect of paeoniflorin on BDNF protein and mRNA expression in hippocampus DG of CUS subjects is illustrated in Fig. 7. Changes to the number of TrkB positive cells in hippocampus DG of subjects were measured by immunohistochemical techniques. Observed within a field of view amplified by ×10-20, TrkB positive cells appeared as granules and their membranes were brownish yellow. ANOVA results suggested that TrkB positive cells significantly differed among groups \( F(3,19) = 4.8, p = 0.033 \). A significant reduction was detected in TrkB positive cells in hippocampus DG of CUS subjects, with significant differences from the control group \( (p = 0.001) \). Compared with the vehicle group, there was a significant increase in the number of TrkB positive cells \( (p = 0.020) \) in hippocampus DG of paeoniflorin treated groups, but no significant difference was obtained for imipramine treated groups \( (p = 0.191) \) (Fig. 7E).

4. Discussion

Extensive studies suggest that most chronic stress inhibits neurogenesis (Dranovsky and Hen, 2006; Sousa et al, 2000; Malberg, 2004). Stress suppresses proliferation of precursor cells in hippocampus DG. It causes morphological changes of neurons and dendritic atrophy of hippocampal pyramidal neurons, thus resulting in DG nerve injuries (Dranovsky and Hen, 2006; Sousa et al, 2000). After modelling with stressors in this experiment, a significant reduction was observed in newborn neural stem cells expressed by BrdU positive cells in hippocampus DG. These results suggest that CUS reduced proliferation of hippocampal neural stem cells in CUS-induced rat with depressive-like behaviors and suppressed neurogenesis.

According to the results, paeoniflorin enhanced the expression of BrdU positive neural cells in the hippocampus of rats, the proliferation of hippocampal neural stem cells and the transformation of hippocampal neural stem cells of CUS rats into astrocytes (the ratio of co-expressed BrdU/GFAP positive cells), but had no significant effect on the transformation of hippocampal neural stem cells into mature neurons in CUS-induced rats. This suggests that paeoniflorin was effective for promoting neurogenesis and transformation of hippocampal neural stem cells into astrocytes. It has previously been reported that astrocytes promote neurogenesis of hippocampal neural stem cells in adult rats (Guillamón-
plasticity in neurons, while their hypofunction might be associ-
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