We investigated the levels of expression of caspase-3 and 9 in the medial prefrontal cortex (mPFC) of rats subjected to single prolonged stress (SPS), trying to provide a novel insight into the mechanism of how this cortical region is related to post-traumatic stress disorder (PTSD). Fifty male Wistar rats were divided into the control group and four SPS groups examined at days 1, 4, 7, and 14 after treatment. Expression of caspase-3 in SPS groups was significantly greater when compared with the control group \((P < 0.05)\) and peaked at day 7 after exposure to SPS. In the control group, the intensity of fluorescence of caspase-9-positive cells was low, while that in the SPS groups was significantly higher \((P < 0.01)\) and peaked at day 4 after exposure to SPS. After SPS episodes, levels of mRNA of caspase-3 and caspase-9, compared with those in the control group, gradually increased and peaked at days 7 and 4, respectively \((P < 0.01)\). Therefore, changes of expression of caspase-9 and caspase-3 may play an important role in the pathogenesis of PTSD.

Keywords: caspase-3, caspase-9, medial prefrontal cortex, single prolonged stress, posttraumatic stress disorder (PTSD).

INTRODUCTION

Post-traumatic stress disorder (PTSD) is an anxiety-related disorder that develops after exposure to a life-threatening traumatic experience. It is characterized by symptoms that often endure for years, including continuous re-experience of the traumatic event, avoidance of stimuli associated with the trauma, numbing of general responsiveness, and increased arousal [1-4]. The pathophysiology of PTSD has been widely studied in neuroscience [5]. However, the mechanism of PTSD is not fully understood until now.

Many lines of evidence from researches on both humans and animals now exist that the amygdala, hippocampus, and medial prefrontal cortex (mPFC) play important roles in PTSD-related symptoms [6]. The amygdala has been documented to play such role in fear, rage, and emotional memory [7, 8]. This cortical zone has direct synaptic connections to amygdala and can inhibit amygdalar activity [9]. The function of mPFC in patients with PTSD undergo negative modifications [10].

In our study, we aimed to explore changes of the caspase 3 and caspase 9 amounts in the mPFC, in order to provide experimental evidence for degeneration of the mPFC in an animal model of PTSD. There are indications that changes in the activities of the above enzymes are important factors influencing the functional state of the mPFC.

The model based on a single prolonged stress (SPS) episode is a good animal model of PTSD related to time-dependent dysregulation of the hypothalamus-pituitary-adrenal axis (HPAA). This model has been developed and employed for PTSD research [11-13]. Many lines of evidence have implicated that shifts in the mPFC play an important role in modulating HPAA responses to emotional stress [14, 15]. Thus, the mPFC deserves special attention in attempts to reveal the mechanism of PTSD.

METHODS

Animals. One hundred healthy male Wistar rats (7 or 8 weeks at the start of the study, weighing approximately 180 g) were supplied for all experiments by the Animal
Experimental Center, China Medical University, were used. All rats were reared in the experimental animal facility for a week to acclimate to new environment (22 ± 1°C, 55 ± 5% humidity, a 12-h light/dark schedule, free access to food and water).

Animal Model and Experimental Groups. The SPS model consisted of 2-h-long whole body restraint in an animal holder. This was followed immediately by a 20-min-long episode of forced swimming (water temperature 25°C, depth of the aquarium 40 cm). The rats were allowed to recuperate for 15 min. Then, they were exposed to inhalation of ether vapors until loss of consciousness. After this, the rats were put back into their home cage and left undisturbed until they were euthanized for the experiments.

The rats were randomly divided into five groups, the normal control group and four SPS-treated groups (1d, 4d, 7d, and 14d). Control rats lived in their individual home cages with no handling and were euthanized after acclimating to their new environment for a week.

Brain Tissue Preparation and Immunohistochemical Analysis of Caspase 3. Rats of each group were transcardially infused with 200-300 ml of cold saline through the ascending aorta followed by 300 ml of 4% cold paraformaldehyde solution. The whole brains were rapidly removed, dissected on ice, and subjected to 6- to 10-h-long post-fixation in 4% paraformaldehyde at 4°C. The brain tissue was cut into 15-μm-thick slices. After treated with 1% hydrogen peroxide/methanol, the sections were incubated with 10% normal goat serum for 30 min at 37°C, with antibody I (rabbit polyclonal anti-caspase 3 antibody, Cell Signaling, USA, dilution 1:300) at 4°C overnight, and with antibody II (goat polyclonal anti-rabbit IgG, Boster Biological Technology, China, dilution 1:200) for 1 h at 37°C. Then, the sections were incubated with avidin-biotin peroxidase complex (Boster Biological Technology, China, dilution 1:200) for 1 h at 37°C. Immunocomplexes were finally visualized with 0.05% diaminobenzidine.

An independent investigator obtained images from five slices for each rat, and the results were analyzed by an image analyzer. The average number of positive cells from the amygdala in each rat (located according to the atlas [16]) was calculated from three sequential brain slices throughout the area of interest and measured for both left and right hemispheres. Counting of positive cells was performed by an individual blind to the treatment conditions, using the same magnification and a identical color scale setting as a correction for background staining.

**Immunofluorescence Analysis of Caspase-9.** The sections were treated with 5% bovine serum albumin (BSA) and 0.3% Triton X-100 in PBS for 30 min at room temperature, to block non-specific staining. Endogenous peroxidase was inactivated with 3% H₂O₂ in double-distilled H₂O for 5 min also at room temperature. The sections were then incubated with mouse anti-caspase 9 monoclonal antibody (Santa Cruz, USA; 1:300) in 2% BSA-PBS overnight at 4°C. After being three times washed with PBS, the sections were incubated with FITC goat anti-mouse IgG (Boster, China; 1:50) for 2 h at room temperature. To assess nonspecific staining, a few sections in every experiment were incubated in the buffer without primary antibody. Slices were then mounted with glycerin and observed using a fluorescence microscope.

Slides were randomly selected from each group. Five visual fields in the mPFC were randomly selected in each slide (×400). The optical density (OD) of caspase-9-immunopositive cells was measured in each field, and the OD average was calculated.

**Reverse Transcription-Polymerase Chain Reaction (RT-PCR) to Detect Caspase-3 and Caspase-9.** Total mRNA of each group was extracted from the basolateral amygdala according to the Trizol kit instructions (Invitrogen, USA), and 1 μg of total RNA was reverse-transcribed into cDNA. The latter was amplified using a RNA PCR kit (AM Ver. 3.0, Takara Bio, Japan). The primers were designed and synthesized by the Shenggong Biotech Company (China) according to the serial number from the Genbank; these data are shown in Table 1. The reaction was started at 94°C for 2 min and amplified.

<table>
<thead>
<tr>
<th>Name</th>
<th>Upstream primer</th>
<th>Downstream primer</th>
<th>Product size(bp)</th>
</tr>
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<tbody>
<tr>
<td>Caspase 3</td>
<td>5'-ctcggtctgtaatgtctgatg-3'</td>
<td>5'-gggtacccggtaggatgtgca-3'</td>
<td>284bp</td>
</tr>
<tr>
<td>Caspase 9</td>
<td>5'-atggaacccgagcccggtctc-3'</td>
<td>5'-ctcattgctctttgagcctcc-3'</td>
<td>330bp</td>
</tr>
<tr>
<td>β-actin</td>
<td>5'-ataacaccaacagtgtgcctc-3'</td>
<td>5'-acagagctgtgcttcagga-3'</td>
<td>542bp</td>
</tr>
</tbody>
</table>

**Table 1. The sequences of caspase 3, caspase 9, and β-actin**
of 40 cycles of 30 sec at 94°C, 60 sec at 52°C (for caspase-3), or 45 sec at 67°C (for caspase-9), 45 sec at 72°C, and ended with a 7-min-long extension at 72°C. β-actin mRNA used as an internal control was co-amplified with caspase 3- or caspase 9-mRNA. The products were observed after electrophoresis on 1.2% agarose gel, and the density of each band was analyzed using the Gel Image Analysis System (Tanon 2500R, China). The levels of caspase 3- and caspase 9-mRNA were estimated by calculating the density ratio of caspase 3 mRNA/β-actin mRNA or caspase 9 mRNA/β-actin mRNA.

Statistical Analysis. All data were expressed as means ± s.d. Intergroups differences were analyzed by one-way analysis of variance (ANOVA) using SPSS 13.0 software. Differences with \( P < 0.05 \) were considered statistically significant.

RESULTS

Immunohistochemical Analysis of Caspase 3. As is shown in Fig. 1A-E, caspase 3 is widely distributed throughout the mPFC region, mainly in the cytoplasm. Labelled structures appeared as Buffy particles. The results indicated that expression of caspase (mean integral optical density) 3 in SPS groups was significantly greater compared to that in the control group \( (P < 0.05) \). This index peaked at day 7 after exposure to SPS (Fig. 1F).

Caspase-9 Immunofluorescence. Immunofluorescence staining results of caspase-9 are shown in Fig. 2. The caspase-9 protein was located in the cytoplasm (Fig. 2A-E). In the normal control group, the fluorescence of caspase-9-positive cells was weak, while that in the SPS groups was significantly

![Fig. 1. Expression of caspase 3 in the medial prefrontal cortex (mPFC) of experimental rats. A) In a rat of the control group; B-E), in rats of the groups subjected to single prolonged stress (SPS) at days (d) 1 (B), 4 (C), 7 (D), and 14 (E). ×400. F) Mean integral optical density, arb. units, in slices from animals of the above groups. *\( P < 0.05 \) vs. the control group.

Рис. 1. Експресія каспази-3 в медіальній префронтальній корі (мPFC) експериментальних щурів.
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**Fig. 2.** Expression of caspase 9 in the prefrontal cortex of experimental rats. A–E) Visualization of expression; F) mean fluorescence intensity, arb. units. Designations are similar to those in Fig. 1.

**Р и с. 2.** Експресія каспази-9 у префронтальній корі експериментальних щурів.

stronger than that in the control ($P < 0.01$). The mean fluorescence intensity in the respective slices peaked at day 4 after exposure to SPS (Fig. 2F).

**RT-PCR Results.** The levels of caspase-3 and caspase-9 mRNA were normalized with respect to the $\beta$-actin mRNA level. The representative gel patterns of caspase-3, caspase-9, and $\beta$-actin cDNA bands are shown in Fig. 3A. After SPS stimulation, caspase-3 and caspase-9 mRNA levels gradually increased, compared with control group, and peaked at days 7 and 4 after stressing, respectively ($P < 0.01$) (Fig. 3B). This result was consistent with that of immunohistochemical analysis.

**DISCUSSION**

PTSD is thought to involve a dysfunction induced by fear-related stimuli. Four major types of typical symptoms of PTSD are the following: re-experiencing, avoidance, numbing, and hyperarousal [1]. Many lines of evidence have implicated a special role of the amygdala; there results were obtained in both animal and human investigations in the processing of threat-related stimuli, in particular anger and fear. These findings are well-documented [2, 17].

Previous studies showed increased activity of the amygdala in patients with PTSD; this was shown by brain imaging. The mPFC can store memories of previous experience, and this plays a key role in the regression on fear conditioning; this cortical area can inhibit amygdalar activity [18]. Studies also showed that declines of mental functions of different degrees in PTSD may be relate to the fact that the mPFC provides descending inhibition of the amygdala inadequately. It was also found that the mPFC volume in PTSD patients decreases [19, 20]. So, dysfunction of the mPFC correlates with reduction of its dimension.
To clarify whether apoptosis is responsible for the decrease in the volume of the mPFC, we estimated the expression of apoptosis-related proteins in the mPFC neurons of rats with PTSD in our study.

Apoptosis is a genetically programmed and morphologically specific form of cell death that can be triggered by a variety of physiological and pathological stimuli. It is well known that caspases, a family of cysteine-dependent aspartate-directed proteases, play critical roles in initiation and execution of apoptosis by cleaving a large number of proteins, which in turn determine typical morphology of apoptosis [3, 21-23]. In the pathogenesis of apoptosis, activation of caspases is a key link, which eventually lead to degradation of chromosomal DNA and cell disintegration. Caspase-9 is upstream of the start caspases, it mainly provides endogenous activation of the apoptotic pathway to activate the effect of caspase-3 immediately leading to apoptosis. We found changes of the levels of caspase-9 and caspase-3 in the mPFC of rats after treatment with single-prolonged stress. The possible reason of the observed pattern is that such stress results in upregulation of the caspases and leads to dysfunction of the mitochondria, which facilitates the apoptotic process. In our study, detection of expressions of caspase-3 and caspase-9 in mPFC neurons, using immunohistochemistry, immunofluorescence, and RT-PCR, demonstrated that both these enzymes are upregulated after stress stimulation, and this provides intensification of apoptosis. A limitation of the current study was that we didn’t examine relations between the mPFC and other involved cerebral structures in order to better explore the mechanism of PTSD.

Thus, we demonstrated that the levels of caspase-3 and caspase-9 significantly increase in the rat mPFC after the animals were subjected to single prolonged stress. The results suggest that the activity of caspase-3 and caspase-9 might play an important role in PTSD. At present, the pathogenesis of PTSD is not yet entirely clear. PTSD may cause a series of biochemical abnormalities and dysfunction of the mPFC, which leads to dysfunctions of other related brain structures. It was obvious that the pathogenesis of PTSD needs to be further studied.

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Experiments were performed in accordance with the National Institute of Health Guide for the Care and Use of Laboratory Animals (NIH Publications No. 80-23) revised 1996. All efforts were made to reduce the number of animals used and to minimize animal suffering during the experiment.

The authors, J. H. Zhang, M. Li, F. Han, and Yu X. Shi, confirm that they have no conflict of interest with any organization or person that may be related to this study; there was also no conflict of interest in interrelations between the authors.
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REFERENCES


