Therapeutic effects of stiripentol against ischemia-reperfusion injury in gerbils focusing on cognitive deficit, neuronal death, astrocyte damage and blood brain barrier leakage in the hippocampus

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ABSTRACT Stiripentol is an anti-epileptic drug for the treating of refractory status epilepticus. It has been reported that stiripentol can attenuate seizure severity and reduce seizure-induced neuronal damage in animal models of epilepsy. The objective of the present study was to investigate effects of post-treatment with stiripentol on cognitive deficit and neuronal damage in the cornu ammonis 1 (CA1) region of the hippocampus proper following transient ischemia in the forebrain of gerbils. To evaluate ischemia-induced cognitive impairments, passive avoidance test and 8-arm radial maze test were performed. It was found that post-treatment with stiripentol at 20 mg/kg, but not 10 or 15 mg/kg, reduced ischemia-induced memory impairment. Transient ischemia-induced neuronal death in the CA1 region was also significantly attenuated only by 20 mg/kg stiripentol treatment after transient ischemia. In addition, 20 mg/kg stiripentol treatment significantly decreased ischemia-induced astrocyte damage and immunoglobulin G leakage. In brief, stiripentol treatment after transient ischemia ameliorated transient ischemia-induced cognitive impairment in gerbils, showing that pyramidal neurons were protected and astrocyte damage and blood brain barrier leakage were significantly attenuated in the hippocampus. Results of this study suggest stiripentol can be developed as a candidate of therapeutic drug for ischemic stroke.

INTRODUCTION

Transient ischemia in the brain can cause metabolic and neurochemical alterations and lead to selective neuronal loss (death) in vulnerable regions after a transient ischemia [1,2]. In particular, neuronal loss occurs in the cornu ammonis (CA) 1 field in the hippocampus proper consisting of CA1 to CA3 fields at a few (four to five) days after five minutes of transient forebrain ischemia (TFI), which is called “delayed neuronal death” [3]. Although the exact mechanism of TFI-induced delayed neuronal death in the hippocampal CA1 field has not been clearly elucidated yet, many previous studies have suggested some underlying mechanisms...
such as excitotoxicity and oxidative stress [4-6]. Neuroinflammatory processes including glial activation and enhanced production and release of inflammatory cytokines have also been thought as major mechanisms [2,7,8]. In addition to delayed neuronal death, it has been demonstrated that TFI can cause the damage of blood-brain barrier (BBB) integrity, leading to leakage and increased permeability (i.e., vasogenic edema, abnormal exchange of molecules, etc.) [9-12].

Stiripentol (STP) is a structurally unique anti-epileptic drug for treating Dravet syndrome and refractory status epilepticus [13]. Previous studies have suggested some underlying mechanisms of the anticonvulsant effect of STP such as action on gamma-aminobutyric acid A receptor and inhibition of lactate dehydrogenase [14,15]. In addition, it has been reported that STP can reduce seizure severity and seizure-induced neuronal damage in animal models of epilepsy [16-18]. In addition, a recent study has reported the effect of STP against neuropathic pain induced by spinal nerve transection in mice, showing that intrathecal administration with STP can attenuate mechanical hyperalgesia in a dose-dependent manner [19].

Although anti-epileptic and neuroprotective effects of STP have been demonstrated in epileptic seizure models, studies investigating whether STP has protective effect against ischemic damage induced by TFI have not been reported yet. Therefore, the objective of this study was to investigate the effects of post-treatment with STP after TFI on TFI-induced cognitive impairment, neuronal damage/death, astrocyte damage and BBB leakage in the hippocampal CA1 region using the gerbil as a good model of TFI [2,20,21].

METHODS

Experimental animals

Male Mongolian gerbils (total n = 144; 80 ± 5 g of body weight; six months of age) were obtained from Experimental Animal Center of Kangwon National University (Chuncheon, Korea). They were housed under optimal conditions (24 ± 0.5°C of room temperature; 55 ± 5% of relative humidity; 12:12 h of light/dark cycle) with three to five animals per cage. Freely accessible pellet feed (DBL Co., Ltd., Eumseong, Korea) and potable water were provided to the gerbils. Entrance to the room was rigorously regulated to reduce superfluous stress to the gerbils.

In this experiment, the experimental protocol was approved (approval no., KW-200113-1) by “Institutional Animal Care and Use Committee” of Kangwon National University on February 18, 2020.

Experimental groups

In this study, experimental groups were as follows: 1) Sham-vehicle group (n = 15) was given sham operation and treated with vehicle (saline); 2) TFI-vehicle group (n = 21) was given TFI operation and treated with vehicle; 3), 4), and 5) Sham-10, 15, and 20 mg/kg STP groups (n = 15, respectively) were given sham operation and treated with 10, 15, and 20 mg/kg STP, respectively; 6), 7), and 8) TFI-10, 15, and 20 mg/kg STP groups (n = 21, respectively) were given TFI operation and treated with 10, 15, and 20 mg/kg STP, respectively.

Surgery for TFI and administration of STP

TFI was induced in accordance with a previously published method [4,12]. In brief, the gerbils were anesthetized by inhaling 2.5% isoflurane (Hana Pharm. Co., Ltd., Seoul, Korea) in 33% O2 and 67% N2O. Under anesthesia, the ventral necks of the gerbils were shaved, and a midline incision (1.2 ± 0.2 cm) was made. Next, bilateral common carotid arteries (BCCA) were isolated and ligated with aneurysm clips (0.69 N) ( Yasargil FE 723K; Aesculap, Tuttingen, Germany). To verify complete occlusion of the BCCA, the central arteries located in bilateral retinas were observed using an ophthalmoscope (HEINE K180) (Heine Optotechnik, Herrsching, Germany). After the conformation of perfect ischemia, the clips were removed at 5 min after BCCA ligation, and the incision was closed using 3-0 suture silk (Ethicon Inc., Somerville, NJ, USA). During the surgical procedure, body temperature was monitored in real-time with a rectal temperature probe (TR-100) (Fine Science Tools, Foster City, CA, USA), and body temperature was controlled at normothermic condition (37 ± 0.5°C). In this experiment, the sham operation was performed like TFI procedure without BCCA ligation.

For the administration of STP (synonyms, 4,4-Dimethyl-1-[(3,4-methylenedioxy)phenyl]-1-penten-3-ol and Diamcon; molecular formula, C14H18O3; solubility, DMSO: ≥ 20 mg/ml), it was dissolved in vehicle (saline) in order that the final dosage was to be 10 mg, 15 mg, and 20 mg per 1 kg of body weight. The gerbils were intraperitoneally injected with vehicle or STP at 30 min after sham or TFI operation (Fig. 1).

Behavioral tests

Passive avoidance test (PAT)

To evaluate short term memory following TFI, PAT was conducted in the light of previous studies [12,22]. In short, we used the Gemini Avoidance System (GEM 392) (San Diego Instruments, San Diego, CA, USA) which consists of dark and light sections communicating by a vertical sliding door between the two sections. As shown in Fig. 1, PAT was performed by two phases at one day before TFI and five days after TFI. Namely, training was performed, and substantive trial was conducted at 20 min after the training. In the training, the gerbil was placed in the light section and allowed to freely explore both sections for one minute while the vertical door was opened. When the gerbil entered the
dark section, the door was closed and an electric foot-shock (0.5 mA for 5 sec) was given to the gerbil from steel grid on the floor. For the substantive trial, the gerbil was placed in the light section, and the latency time to enter the dark section was recorded within three minutes.

**8-arm radial maze test (8-ARMT)**

8-ARMT was performed in order to evaluate spatial memory following TFI in accordance with previously described methods [4,11,12]. Briefly, a central platform consisting of a non-transparent acryl board (diameter, 20 cm) with radially extended eight arms (width, 5 cm; height, 9 cm; length, 35 cm) (Stoelting Co., Wood Dale, IL, USA) was used. Pelleted diets (DBL Co., Ltd.) were located at the end of each arm. As shown in Fig. 1, the gerbils were pre-trained once a day for three days before TFI induction. The substantive trials were conducted once a day for five days from one day after TFI induction. To evaluate 8-ARMT, the gerbil was placed onto the central platform, and the numbers of errors were recorded at every occasion when the gerbil went into the arm that they had already entered before. When the gerbil ate up the pellets, each trial was terminated.

**Preparation of histological sections**

As shown in Fig. 1, the gerbils used for all groups were sacrificed (n = 5, respectively, in the sham groups; n = 7, respectively, in the TFI groups) at six hours, two days and five days after sham or TFI operation. According to our previous studies [4,12], brain tissue sections containing the hippocampus were prepared. In short, the gerbils were profoundly anesthetized by intraperitoneal administration of 90 mg/kg pentobarbital sodium (JW pharm. Co., Ltd., Seoul, Korea). Under deep anesthesia, the brains of the gerbils were rinsed with saline through ascending aorta and fixed 4% paraformaldehyde (in 0.1 M phosphate buffer; pH 7.4). The brains were harvested and more fixed with same fixative for four hours at room temperature. To cryoprotect the brains when they were sectioned, the fixed brains were infiltrated with 25% sucrose solution (in 0.1 M PB; pH 7.4) for 24 h at room temperature. Using a sliding microtome (SM2020 R) (Leica, Nussloch, Germany) equipped with BFS-40MP freezing platform (Physitemp Instruments, Inc., Clifton, NJ, USA), the brains containing the hippocampus were coronally sectioned into 30 μm of thickness. The sections used in this experiment were selected at level of –1.8 mm to 2.7 mm of antero-posterior to the bregma in the light of the gerbil brain atlas [23].

**Immunohistochemistry**

Immunohistochemical studies were done to examine changes in neurons, astrocytes, and immunoglobulin G (IgG) using the following primary antibodies: neuronal nuclei (NeuN; a marker for neurons), and IgG (Table 1). In brief, in accordance with previously published methods [4,12], the sections were incubated in 0.3% H₂O₂ for 25 min at room temperature, and they were immersed in five percent normal horse or goat serum for 30 min at room temperature. Thereafter the sections were immunoreacted with each primary antibody for 24 h at 4°C. After washing them,
the immunoreacted sections were reacted with corresponding secondary antibodies for two hours at room temperature followed by incubated in diluted avidin-biotin complex (dilute rate, 1:250) (Vector Laboratories, Burlingame, CA, USA) (Table 1). After washing, these sections were subsequently visualized via immersing into 0.06% 3,3’-diaminobenzidine tetrahydrochloride (Sigma-Aldrich Co., St. Louis, MO, USA) containing 0.003% H₂O₂. Finally, these sections were mounted onto the slide glass, dehydrated using ethyl alcohol series, cleared in xylene, and coverslipped with Canada balsam (Kanto Chemical, Tokyo, Japan).

To evaluate the change of NeuN immunoreactive cells (neurons) was performed according to published papers [4,24] with minor modification. The image of NeuN immunoreactive neurons was taken at the magnification of 20X using light microscope (BX53) (Olympus, Tokyo, Japan) equipped with digital camera (DP72) (Olympus). NeuN immunoreactive neurons were counted using image capture software (cellSens Standard) (Olympus). The total number of NeuN immunoreactive neurons was counted in 250 μm² at the middle in the CA1 region, and the mean number was calculated using NIH Image (1.59 software) (National Institutes of Health, Bethesda, MD, USA).

To quantitatively analyze changes in the immunoreactivities of glial fibrillary acidic protein (GFAP), and IgG in the CA1 region, immunostained sections were observed in the same way described above. To evaluate changes in the immunoreactivities, in accordance with previous studies [12,25], all immunoreactive structures were converted into 8-bit grey scale images with a range of 0-255 (from black to white). Using Image J software (version 1.46) (National Institutes of Health), images were assessed for grey scale intensity, and average immunoreactive intensity in each immunoreactive staining was calculated within the area of interest. The immunoreactive intensity was relatively presented as relative optical density (ROD): the optical density of the Sham vehicle group was considered as 100%.

**Histofluorescence using Fluoro-Jade B (F-J B)**

F-J B is used as a marker of cellular degeneration. F-J B histofluorescence was carried out in order to examine neuronal damage/death following TFI. As described previously [12,26], in brief, the sections were incubated in 0.06% potassium permanganate (KMnO₄) (Sigma-Aldrich Co.) for 20 min on an orbital shaker. After washing for two minutes, these sections were reacted in 0.0004% F-J B (Histochem, Jefferson, AR, USA) for 30 min. Thereafter, the brain sections were washed three times for two minutes and placed on a slide warmer until they were fully dried. The slides were cleared using xylene for one minute and coverslipped with dibutylphthalate polystyrene xylene (Sigma-Aldrich Co.).

The count of F-J B positive cells (neurons) was performed to evaluate TFI-induced neuronal death in the CA1 region. In short, according to published methods [4,24], the image of F-J B positive cells was taken at the magnification of 20X using epifluorescent microscope (Carl Zeiss, Oberkochen, Germany), equipped with a digital camera (DP72) (Olympus), at 450–490 nm of wavelength. F-J B positive cells were taken using image capture software (cellSens Standard) (Olympus). The total number of F-J B positive cells was counted in the same way described above (count of NeuN immunoreactive neurons).

**Double immunofluorescence staining**

To distinguish astrocyte endfeet and blood vessels, double immunofluorescence staining was conducted according to a published protocol [12]. In short, the sections were reacted in mouse anti-GFAP (dilute rate, 1:800) (Chemicon, Temecula, CA, USA), mouse anti-NeuN (dilute rate, 1:800) (Chemicon), rabbit anti-glucose transporter 1 (GLUT-1; a marker for endothelial cells) (dilute rate 1:1,000) (Chemicon), and rabbit anti-IgG (dilute rate, 1:800) (Bioss antibodies, Atlanta, GA, USA) were used as primary antibodies. Thereafter, the sections were immunoreacted with secondary antibodies combined Alexa Fluor 488-conjugated donkey anti-mouse IgG (dilute rate, 1:500) (Invitrogen, Waltham, MA, USA) and Alexa Fluor 546-conjugated goat anti-rabbit IgG (dilute rate 1:500) (Invitrogen). After washing, the sections were mounted onto the slide glasses and dehydrated using a dry oven (WiseVen WOC High Clean Air Oven) (Daihan Scientific Co., Ltd., Wonju, Korea) for eight hours. Finally, the sections were coverslipped with DPX (Sigma-Aldrich Co.).

The immunoreaction of GFAP/GLUT-1 was observed using confocal MS (LSM510 META NLO) (Carl Zeiss), which was in the Korea Basic Science Institute Chuncheon Center (Chuncheon, Korea).

**Statistical analysis**

In this study, we used SPSS software (version 15.0) (SPSS Inc., Chicago, IL, USA) to perform statistical analyses. Kolmogorov and Smirnov test was used to test normal distributions, and Bartlett test was used to test identical standard error of the means (SEMs). All our data passed the normality test. The statistical significances of the mean among the experimental groups were determined by two-way analysis of variance (for the numbers of NeuN-immunoreactive neurons and F-J B positive cells, RODs of GFAP and IgG-immunoreactive structures, and latency times in PAT) and repeated measures analysis of variance (for the numbers of errors in 8-ARMT) followed by post-hoc Tukey test for all pairwise multiple comparisons. Statistical significance was considered to be at a value of p < 0.05.
RESULTS

Effect of STP against TFI-induced cognitive deficit

In this study, as shown in the Fig. 2, STP treatment after TFI significantly attenuated TFI-induced cognitive impairments. In PAT, the latency time one day before TFI was not significantly different between the Sham-vehicle and all Sham-STP groups, and, five days after TFI, the latency time in all Sham groups was not different from that shown one day before TFI (Fig. 2A). In the TFI-vehicle, TFI-10 mg/kg STP and TFI-15 mg/kg STP groups, the latency time five days after TFI was significantly lower than that in the Sham-vehicle and Sham-STP groups (Fig. 2A). However, five days after TFI, the latency time in the TFI-20 mg/kg STP group was similar to that found in the Sham-vehicle group (Fig. 2A).

In 8-ARMT, the number of the errors was gradually decreased in all groups with time before TFI, and the decrease pattern was not different between all groups (Fig. 2B). One day after TFI, the numbers of the errors were significantly increased in the TFI-vehicle and all TFI-STP groups when compared with the Sham-vehicle and all Sham-STP groups (Fig. 2B). From two days to five days after TFI, the numbers of the errors in the TFI-vehicle, TFI-10 mg/kg STP and 15 mg/kg STP groups were not significantly decreased with time after TFI (Fig. 2B). However, the numbers of the errors in the TFI-20 mg/kg STP group began to decrease from two days after TFI and were significantly decreased with time, showing that the numbers five days after TFI were similar to those shown in the Sham groups (Fig. 2B).

Effect of STP against TFI-induced neuronal death

In this experiment, we found that STP treatment after TFI protected neurons in the CA1 region from TFI injury (Fig. 3). In the Sham-vehicle group, neurons immunostained with NeuN (NeuN- neurons), as intact neurons, were located in the stratum pyramidale, and cells positive to F-J B (F-J B’ cells), as damaged (dead) cells, were not found (Fig. 3A, 3a, 3I, 3i). On the other hand, in the TFI-vehicle group, very few NeuN- neurons and many F-J B’ cells were shown in the stratum pyramidale five days after TFI (Fig. 3B, 3b, 3I, 3i).

In all Sham-STP groups, no significant differences in the numbers of NeuN- and F-J B’ cells were shown when compared with those found in the Sham-vehicle group (Fig. 3C, 3c, 3E, 3e, 3G, 3g, 3I, 3i). In the TFI-10 mg/kg STP and TFI-15 mg/kg STP groups, the distribution of NeuN- and F-J B’ cells was very similar to the TFI-vehicle group (Fig. 3D, 3d, 3F, 3f, 3I and 3i). In contrast, in the TFI-20 mg/kg STP group, many NeuN- and very few F-J B’ cells were detected in the stratum pyramidale (Fig. 3H, 3h, 3I, 3i): this finding means that therapeutic treatment with 20 mg/kg STP after TFI in gerbils very effectively protected hippocampal neurons from ischemia-reperfusion injury.

Based on these results, we described the following results obtained from the TFI-20 mg/kg STP group.

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Fig. 2. Latency time in PAT (A) and mean numbers of errors in 8-ARMT (B) in the Sham-vehicle, Sham-STP (10, 15, and 20 mg/kg), TFI-vehicle and TFI-STP (10, 15, and 20 mg/kg) groups before and after TFI. The substantial trials for PAT are conducted one day before and five days after TFI and gerbils are undergone pre-training trial at 20 min before each substantial trial. In the TFI-20 mg/kg STP group, a significantly delayed latency time in PAT is recorded compared with that in the TFI-vehicle, TFI-10 mg/kg STP and TFI-15 mg/kg STP groups. 8-ARMT is daily performed at three to one day before TFI, and one to five days after TFI. In the TFI-20 mg/kg STP group, the numbers of the errors in 8-ARMT are significantly decreased from two days after TFI. The bars indicate the means ± SEM (n = 7, respectively; *p < 0.05 vs. Sham-vehicle or Sham-STP group; #p < 0.05 vs. TFI-vehicle group). 8-ARMT, 8-arm radial maze test; PAT, passive avoidance test; STP, stiripentol; TFI, transient forebrain ischemia.
Effect of STP against TFI-induced astrocyte damage

We found, in this study, that STP treatment after TFI significantly attenuated TFI-induced astrocyte damage in the CA1 region (Fig. 4). In the Sham-vehicle group, cells immunostained with GFAP, as intact GFAP⁺ astrocytes, had thin processes and small cytoplasm, and most of them were distributed in stratum oriens and radiatum (Fig. 4Aa). In the TFI-vehicle group, GFAP immunoreactivity in the astrocytes was increased with time after TFI, showing that their cytoplasm and processes became gradually hypertrophied and thickened, respectively (Fig. 4Ab–Ad and 4B).

No significant difference was shown in GFAP immunoreactivity between the Sham-20 mg/kg STP group and the Sham-vehicle groups (Fig. 4Ae, 4B). In the TFI-20 mg/kg STP group, the hypertrophy of the GFAP⁺ astrocytes was significantly attenuated after TFI when compared with those found in the Sham-vehicle group (Fig. 4Af–Ah and 4B).

To examine the changes of the ends of astrocyte processes, as astrocyte endfeet (AEF), we performed double immunofluorescence for GFAP and GLUT-1 in the CA1 region after TFI. In all Sham groups, GFAP⁺ AEF were found around GLUT-1⁺ endothelial cells (Fig. 5A, E). In the TFI-vehicle group, the AEF disappeared gradually after TFI, showing that, five days after TFI, most of GFAP⁺ AEF were not contact with small blood vessels (Fig. 5B–D). On the other hand, the damage of GFAP⁺ AEF was significantly attenuated in the TFI-20 mg/kg STP-group (Fig. 5F–H).

Effect of STP against TFI-induced IgG leakage

As shown in the Fig. 6, in this study, we found that STP treatment after TFI significantly attenuated TFI-induced IgG leakage in the CA1 region. In the Sham-vehicle group, IgG immunoreactivity was shown inside blood vessels, not in CA1 parenchyma (Fig. 6Aa). In the TFI-vehicle group, IgG immunoreactivity was shown in cells six hours after TFI, apparently increased in the parenchyma and cells two days after TFI, and very high five days after TFI (Fig. 6Ab–Ad and 6B).

In the Sham-20 mg/kg STP group, IgG immunoreactivity was not different from the Sham-vehicle group (Fig. 6Ae). In the TFI-20 mg/kg STP group, IgG immunoreactivity was significantly lower than that in the TFI-vehicle group, although the distribution pattern of IgG⁺ structures was similar to that shown in the TFI-vehicle group (Fig. 6Af–Ah and 6B).
DISCUSSION

In this study, to investigate the therapeutic effect of STP against ischemic injury induced by TFI and its possible mechanism, cognitive impairment, neuronal death, astrocyte damage, IgG leakage and microgliosis were examined in the CA1 region of gerbil
hippocampus known to be vulnerable to transient ischemia, following 5-min TFI.

It is well known that TFI-induced hippocampal neuronal damage is closely associated with deficits in learning and memory and that impaired cognitive performance can occur before the onset of TFI-induced neuronal loss (death) in the hippocampus [10-12,27]. PAT and 8-ARMT have been widely used to investigate alterations of cognitive function following ischemic injury in the hippocampus induced by ischemia-reperfusion [11,12,22,28]. Therefore, in the present study, to examine whether STP treatment after TFI in gerbils affected TFI-induced change in cognitive function, we performed PAT and 8-ARMT. Result of PAT revealed that the latency time in the TFI-20 mg/kg STP group was very similar to that in the Sham-vehicle group five days after TFI. Result of 8-ARMT showed that the number of errors in the TFI-20 mg/kg STP group was not significantly different from that in the Sham-vehicle group. These results indicate that 20 mg/kg STP (not 10 and 15 STP) treatment after TFI can improve TFI-induced cognitive impairment.

Some previous studies have reported the neuroprotective effect of STP in experimental animal models of neurological disorders. Auvin et al. [16] reported that STP reduced neuronal injury in the hippocampus of a rat model of status epilepticus induced by lithium-pilocarpine. In particular, Verleye et al. [29] used two in vitro models of brain injury associated with seizure through neuronal-astroglial cultures from rat cerebral cortex exposed to high level of glutamate (40 µM for 20 min) or to oxygen-glucose deprivation (OGD) and found that STP had neuroprotective effects when it was used for treatment before, but not after, exposure to GOD, whereas neuroprotective effects by STP was found when it was treated both before and after exposure to a high level of glutamate. In addition, Chatterjee et al. [30] recently reported that STP had a neuroprotective effect against neurotoxicity in PC12 cells induced by treatment with anti-nerve growth factor and that STP treatment reduced neurotoxicity in rat primary cortical neurons induced by Aβ42 implicated in Alzheimer's disease. In the present study, we examined neuroprotective effects of post-treatment with STP against TFI-induced ischemic damage in gerbil hippocampal CA1 region using NeuN immunohistochemical staining and F-J B histofluorescence, and found that post-treatment with STP at 20 mg/kg, but not at 10 or 15 mg/kg, protected against TFI-induced delayed neuronal death in the hippocampal CA1 region. Based on our results and previous findings, to the best of our knowledge, this is the first study showing that STP had a neuroprotective effect against ischemia-reperfusion injury using a gerbil model of TFI. This result is correlated with results of the two behavioral tests (PAT and 8-ARMT).

Astrocytes (astroglial cells) are the most common glial cells located in the central nervous system (CNS; brain and spinal cord). They perform diverse functions including maintenance of extracellular ion balance, support of endothelial cells for the constitution of BBB, nutrient provision to nervous tissues, and participation in repair of CNS injuries [31-33]. It has been reported that astrocytes play significant roles in both healthy and
injured brains including ischemic stroke [34-37]. Functional and morphological alterations in astrocytes have been reported in brains with pathological conditions (the process is termed "astroglisis") [2,38,39]. It is known that astroglisis is characterized by proliferation and hypertrophy of astrocytes with increased GFAP expression in astrocytes [2,40]. However, Sofroniew and Vinters [41] have reported that an increase of cells positive to GFAP does not result from new astrocyte generation, but from an increase in GFAP synthesis and filament condensation in pre-existing astrocytes. Based on this finding, we have recently reported that GFAP positive cells (astrocytes) with strong GFAP immunoreactivity in gerbil striatum are swollen in cell bodies and destroyed (short and thickened) in their processes at five days after TFI [39]. In the current study, GFAP immunoreactivity was significantly increased in astrocytes of the TFI-vehicle group. Their cytoplasm and processes became gradually hypertrophied and thickened. Furthermore, when we performed double immunofluorescence for GFAP and GLUT-1, AFE (ends of astrocyte processes) disappeared seriously at five days after TFI. Most of the GFAP positive AFE were not having contacts with small blood vessels. However, in the TFI-20 mg/kg STP group, the hypertrophy of GFAP positive astrocytes was significantly attenuated after TFI. Furthermore, the damage of AEF was significantly attenuated. For reference on AEF, Abbott [42] reported that the AEF of astrocytes envelopes almost all (> 99%) endothelium in brains. In addition, Daneman and Prat [32] have reported that astrocytes provide cellular linkages between neurons and small blood vessels by AFE which is a part of the BBB [2]. Taken together, our results suggest that the damage of astrocytes (in particular, AFE) in the hippocampus after TFI is involved in TFI-induced neuronal loss (death), which is related to cognitive deficit, and that this damage can be attenuated or protected by 20 mg/kg STP.

Finally, we investigated whether post-treatment with 20 mg/kg STP displayed a protective effect against TFI-induced BBB leakage. A healthy condition in the CNS is attained by BBB through preserving the balance between water and ion and supplying nutrients [31,32]. In addition, it is generally accepted that BBB integrity should be maintained to protect brain following an ischemic injury. Thus, maintenance of the BBB integrity is considered as one of potential therapeutic targets for treating an ischemic stroke [43-45]. In normal conditions, BBB as a selectively permeable barrier allows only specific substances from blood to enter brain parenchyma and participate in maintaining homeostasis in the CNS. However, disruption of BBB integrity, BBB leakage and marked extravasation can occur after an ischemic injury [10,12,46,47]. In many experiments, extravascular IgG has been used to examine the increase of BBB permeability [10,12,47]. Therefore, we performed IgG immunohistochemistry and found that IgG immunoreactivity was markedly increased in the CA1 region after TFI. We then we investigated the protective effect of STP against TFI-induced BBB leakage. We found that STP treatment significantly decreased IgG immunoreactivity in the ischemic CA1 region. This finding indicates that STP treatment after TFI can protect the BBB integrity in ischemic brains.

In conclusion, this study showed that post-treatment with 20 mg/kg STP ameliorated TFI-induced cognitive impairment and protected against TFI-induced neuronal death in the hippocampal CA1 region. In addition, STP treatment markedly attenuated IgG leakage in the ischemic CA1 region. Results of this study suggest that STP can be developed as a therapeutic drug for treating ischemia-reperfusion injury or ischemic stroke.

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**CONFLICTS OF INTEREST**

The authors declare no conflicts of interest.

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