**Scutellaria baicalensis** stem-leaf total flavonoid reduces neuronal apoptosis induced by amyloid beta-peptide (25–35)☆

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**Abstract**

*Scutellaria baicalensis* stem-leaf total flavonoid might attenuate learning/memory impairment and neuronal loss in rats induced by amyloid beta-peptide. This study aimed to explore the effects of *Scutellaria baicalensis* stem-leaf total flavonoid on amyloid beta-peptide-induced neuronal apoptosis and the expression of apoptosis-related proteins in the rat hippocampus. Male Wistar rats were given intragastric administration of *Scutellaria baicalensis* stem-leaf total flavonoid, 50 or 100 mg/kg, once per day. On day 8 after administration, 10 μg amyloid beta-peptide (25–35) was injected into the bilateral hippocampus of rats to induce neuronal apoptosis. On day 20, hippocampal tissue was harvested and probed with the terminal deoxyribonucleotidyl transferase-mediated biotin-16-dUTP nick-end labeling assay. *Scutellaria baicalensis* stem-leaf total flavonoid at 50 and 100 mg/kg reduced neuronal apoptosis induced by amyloid beta-peptide (25–35) in the rat hippocampus. Immunohistochemistry and western blot assay revealed that expression of the pro-apoptotic protein Bax, cytochrome c and caspase-3 was significantly diminished by 50 and 100 mg/kg *Scutellaria baicalensis* stem-leaf total flavonoid, while expression of the anti-apoptotic protein Bcl-2 was increased. Moreover, 100 mg/kg *Scutellaria baicalensis* stem-leaf total flavonoid had a more dramatic effect than the lower dosage. These experimental findings indicate that *Scutellaria baicalensis* stem-leaf total flavonoid dose-dependently attenuates neuronal apoptosis induced by amyloid beta-peptide in the hippocampus, and it might mediate this by regulating the expression of Bax, cytochrome c, caspase-3 and Bcl-2.

**Key Words**

neural regeneration; traditional Chinese medicine; neurodegenerative disease; *Scutellaria baicalensis* stem-leaf total flavonoid; amyloid beta-peptide; neurons; apoptotic protein; cytochrome c; Alzheimer's disease; grants-supported paper; neuroregeneration

**Research Highlights**

(1) Based on the objective criteria for drug effects described in previous in vivo studies, apoptosis was examined in this study, and the expression of apoptosis-related proteins was qualitatively and semi-quantitatively assayed using immunohistochemistry. Furthermore, western blot analysis was used for quantitating protein expression. Bax, Bcl-2, cytochrome c and caspase-3 were selected for assessment.

(2) *Scutellaria baicalensis* stem-leaf total flavonoid can attenuate apoptosis in the hippocampus induced by amyloid beta-peptide, reduce the expression of Bax, cytochrome c and caspase-3, and increase the expression of the anti-apoptotic protein Bcl-2.

(3) *Scutellaria baicalensis* stem-leaf total flavonoid has inhibitory effects against neuronal apoptosis in amyloid beta-peptide (25–35)-treated rats, and the effect is dose-dependent.
INTRODUCTION

Alzheimer’s disease is a degenerative disease of the central nervous system, characterized by progressive cognitive dysfunction and memory impairment. Amyloid beta-protein is widely regarded as the main culprit in the pathogenesis of Alzheimer’s disease. Intracellular accumulation of amyloid beta oligomer may trigger endoplasmic reticulum stress-induced apoptosis. Amyloid beta-peptide (25–35) can induce PC12 cell apoptosis and caspase-dependent and -independent apoptosis of human embryonic neurons.

Scutellaria baicalensis stem-leaf total flavonoid is extracted from the stems and leaves of Scutellaria baicalensis Georgi. Scutellaria baicalensis stem-leaf total flavonoid has anti-inflammatory and immune modulatory effects, and can alleviate myocardial ischemia-reperfusion injury. In addition, the antioxidative effects of Scutellaria baicalensis stem-leaf total flavonoid help to attenuate learning and memory impairment, and prevent neuronal loss and ultrastructural damage in the hippocampal CA1 region after injection of amyloid beta. 5,7-Dihydroxy flavanone might inhibit mitochondrial-mediated apoptosis and regulate the expression of apoptosis-related proteins after intraventricular injection of amyloid beta-peptide (25–35) into mice. Curcumin was shown to mediate neuroprotection against the mitochondrial apoptosis pathway. However, there is little research on the effects of Scutellaria baicalensis stem-leaf total flavonoid on apoptosis.

In this study, we sought to explore the effects of Scutellaria baicalensis stem-leaf total flavonoid on amyloid beta-induced neuronal apoptosis in the hippocampus. We also examined its effects on the expression of the apoptosis-related proteins Bax, Bcl-2, caspase-3 and cytochrome c using the terminal deoxynucleotidyl transferase-mediated biotin-16-dUTP nick-end labeling (TUNEL) assay, immunohistochemistry, and western blot analysis.

RESULTS

Quantitative analysis of experimental animals

Forty male Wistar rats were randomly divided into a control group (intragastric administration of saline + hippocampal injection of sterile saline), model group [intragastric administration of saline + hippocampal injection of amyloid beta-peptide (25–35)], high-dose treatment group [intragastric administration of 100 mg/kg Scutellaria baicalensis stem-leaf total flavonoid + hippocampal injection of amyloid beta-peptide (25–35)], and low-dose treatment group [intragastric administration of 50 mg/kg Scutellaria baicalensis stem-leaf total flavonoid + hippocampal injection of amyloid beta-peptide (25–35)]. All 40 rats were involved in the final analysis.

Scutellaria baicalensis stem-leaf total flavonoid inhibited amyloid beta-peptide (25–35)-induced apoptosis in the rat hippocampus

TUNEL staining revealed numerous brown nuclei in the hippocampus of rats in the model group, while only a few stained cells were found in the control group. Compared with the model group, hippocampal cells were lightly stained in the high-dose and low-dose treatment groups, and there were fewer brown cells (Figure 1).

![Figure 1](image)

**Figure 1** Effect of Scutellaria baicalensis stem-leaf total flavonoid on cell apoptosis (arrows) in the hippocampal CA1 region of amyloid beta-peptide (25–35)-treated rats (terminal deoxynucleotidyl transferase-mediated biotin-16-dUTP nick-end labeling, × 400).

(A) Control group: Cells are arranged in neat rows; a few cells exhibit light yellow nuclei.

(B) Model group: Cells are arranged in a disordered manner; a large number of cells exhibit deeply stained nuclei.

(C) Low-dose Scutellaria baicalensis stem-leaf total flavonoid group: Compared with the model group, cells are arranged tightly, the number of yellow stained cells is reduced, and the staining is light.

(D) High-dose Scutellaria baicalensis stem-leaf total flavonoid group: Compared with the model group, cells are arranged tightly, the number of yellow stained cells is significantly reduced, and the staining is light.

Statistical analysis showed that the number of TUNEL-positive cells in the model group was significantly higher than in the control group (P < 0.01). Furthermore, 50 and 100 mg/kg Scutellaria baicalensis stem-leaf total flavonoid reduced the number of TUNEL-positive hippocampal cells in animals.
administered amyloid beta-peptide (25–35). This effect was more evident at 100 mg/kg ($P < 0.01$; Figure 2).

\[ \text{Figure 2} \quad \text{Effect of Scutellaria baicalensis stem-leaf total flavonoid (SSTF) on cell apoptosis in the hippocampal CA1 region of amyloid beta-peptide (25–35)-treated rats.} \]

Data are expressed as mean ± SD, there were two rats in each group, and four slices ($\times 400$) were selected from each rat. $^aP < 0.01$, vs. control group; $^bP < 0.01$, vs. model group (one-way analysis of variance, least significant difference between two groups).

TUNEL: Terminal deoxyribonucleotidyl transferase-mediated biotin-16-dUTP nick-end labeling.

Scutellaria baicalensis stem-leaf total flavonoid inhibited Bax expression in the hippocampus of amyloid beta-peptide (25–35)-treated rats

Immunohistochemical staining demonstrated the presence of Bax-positive cells exhibiting a brown granular cytoplasm. In the model group, most of the neurons in the hippocampal CA1 region were stained yellow. In contrast, Bax-positive cells in the low and high-dose treatment groups were lightly stained compared with the model group (Figure 3). Statistical analysis showed that Bax expression in the hippocampal CA1 region in the model group was significantly higher than in the control group ($P < 0.01$). Low- and high-dose Scutellaria baicalensis stem-leaf total flavonoid treatment decreased Bax expression in the hippocampal CA1 region ($P < 0.01$), especially the high dose ($P < 0.01$; Table 1).

Western blot analysis results were consistent with the immunohistochemical assay findings. Amyloid beta-peptide (25-35) up-regulated Bax expression, while Scutellaria baicalensis stem-leaf total flavonoid down-regulated Bax expression in the rat hippocampus ($P < 0.01$). The 100 mg/kg dose had a particularly dramatic effect ($P < 0.01$; Figure 4, Table 1).

\[ \text{Figure 3} \quad \text{Effect of Scutellaria baicalensis stem-leaf total flavonoid on Bax expression (arrows) in the hippocampal CA1 region of amyloid beta-peptide (25–35)-treated rats (immunohistochemical staining, $\times 400$).} \]

(A) Control group: Cells are arranged in neat rows, and a few cells have lightly stained cytoplasm.

(B) Model group: Cells are arranged in a disorderly manner, cell count is reduced, and many cells have deeply stained cytoplasm.

(C) Low-dose Scutellaria baicalensis stem-leaf total flavonoid group: Compared with the model group, cells are arranged tightly, the number of stained cells is reduced, and the staining is light.

(D) High-dose Scutellaria baicalensis stem-leaf total flavonoid group: Compared with the model group, cells are arranged tightly, cell count is higher, the number of stained cells is significantly reduced, and the staining level is light.

Scutellaria baicalensis stem-leaf total flavonoid enhanced Bcl-2 expression in the hippocampus of amyloid beta-peptide (25–35)-treated rats

Immunohistochemistry showed that Bcl-2 was mainly expressed in the cytoplasm of rat hippocampal neurons, and was displayed as brownish yellow staining. The number of Bcl-2-positive cells in the hippocampal CA1 region in the model group was less than in the control group, with light staining. In comparison, Bcl-2-positive cells in the low- and high-dose treatment groups were more numerous than in the model group, with intense staining (Figure 5).

\[ \text{Table 1} \quad \text{Effect of Scutellaria baicalensis stem-leaf total flavonoid (SSTF) on Bax expression in the hippocampal CA1 region of amyloid beta-peptide (25–35)-treated rats} \]

| Group      | Immunohistochemical staining (absorbance) | Western blot assay (absorbance ratio of Bax to $\beta$-actin) |
|------------|-------------------------------------------|-------------------------------------------------------------|
| Control    | $20.78 \pm 0.17$                          | $0.10 \pm 0.02$                                             |
| Model      | $144.51 \pm 30.45^{a}$                    | $0.27 \pm 0.05^{a}$                                        |
| Low-dose SSTF | $87.65 \pm 24.56^{b}$                     | $0.20 \pm 0.04^{b}$                                        |
| High-dose SSTF | $52.17 \pm 12.78^{b}$                    | $0.16 \pm 0.03^{b}$                                        |

Data are expressed as mean ± SD. There were two rats in each group, and four slices ($\times 400$) were selected from each for immunohistochemistry. Six rats were taken from each group for western blot analysis. $^{a}P < 0.01$, vs. control group; $^{b}P < 0.01$, vs. model group (one-way analysis of variance, least significant difference between two groups).
Western blot effect (Scutellaria baicalensis) increased significantly after low-dose SSTF (P < 0.01). Furthermore, compared with the model group, the number of deeply stained cells is significantly increased. Statistical analysis showed that, compared with the control group, Bcl-2 expression in the hippocampal CA1 region in the model group was significantly lower (P < 0.01). Furthermore, compared with the model group, Bcl-2 expression in the hippocampal CA1 region was increased significantly after low- and high-dose Scutellaria baicalensis stem-leaf total flavonoid treatment (P < 0.01). High-dose treatment had a more dramatic effect (P < 0.01; Table 2).

Western blot analysis results were consistent with the immunohistochemical results. Amyloid beta-peptide (25–35) down-regulated hippocampal Bcl-2 expression (P < 0.01), and 100 mg/kg Scutellaria baicalensis stem-leaf total flavonoid up-regulated Bcl-2 expression in the rat hippocampus (P < 0.05; Figure 6, Table 2).

Table 2  Effect of Scutellaria baicalensis stem-leaf total flavonoid (SSTF) on Bcl-2 expression in the hippocampal CA1 region of amyloid beta-peptide (25–35)-treated rats

| Group   | Immunohistochemical staining (absorbance) | Western blot assay (absorbance ratio of Bcl-2 to β-actin) |
|---------|------------------------------------------|--------------------------------------------------------|
| Control | 115.24±24.56                             | 0.41±0.07                                             |
| Model   | 21.45±5.27c                              | 0.24±0.04c                                            |
| Low-dose SSTF | 46.27±8.37h                          | 0.24±0.03                                            |
| High-dose SSTF | 76.74±18.19b                           | 0.30±0.04c                                            |

Data are expressed as mean ± SD. There were two rats in each group, and four slices (× 400) were selected from each rat for immunohistochemistry. Six rats in each group were taken for western blot analysis. aP < 0.01, vs. control group; bP < 0.01, cP < 0.05, vs. model group (one-way analysis of variance, least significant difference between two groups).

Scutellaria baicalensis stem-leaf total flavonoid inhibited caspase-3 expression in the hippocampus of amyloid beta-peptide (25–35)-treated rats

Immunohistochemical staining showed that caspase-3 was expressed in the cytoplasm and nuclei of rat hippocampal neurons. The caspase-3-positive cells in the hippocampal CA1 region of rats in the model group were more numerous than in the control group and were deeply stained. Compared with the model group, the number of caspase-3-positive cells was decreased after low- and high-dose Scutellaria baicalensis stem-leaf total flavonoid treatment, and these cells were slightly stained (Figure 7). Statistical analysis revealed that caspase-3 expression in the hippocampal CA1 region in the model group was increased significantly compared with the control group (P < 0.01). Compared with the model group, caspase-3 expression was significantly reduced after low- and high-dose Scutellaria baicalensis stem-leaf total flavonoid treatment.
flavonoid treatment \( (P < 0.01) \), especially with high-dose treatment \( (P < 0.01; \text{Table 3}) \).

![Figure 7](image)

**Figure 7** Effect of *Scutellaria baicalensis* stem-leaf total flavonoid on caspase-3 expression (arrows) in the hippocampal CA1 region of amyloid beta-peptide (25–35)-treated rats (immunohistochemical staining, \( \times 400 \)).

(A) Control group: Cells are arranged in neat rows, and a few cells are stained lightly.

(B) Model group: Cells are arranged in a disorderly manner, and the number of deeply stained cells is increased.

(C) Low-dose *Scutellaria baicalensis* stem-leaf total flavonoid group: Compared with the model group, cells are arranged in neat rows, the number of stained cells is decreased, and staining intensity is low.

(D) High-dose *Scutellaria baicalensis* stem-leaf total flavonoid group: Compared with the model group, the number of stained cells is significantly decreased, and staining intensity is low.

Western blot analysis results were consistent with the immunohistochemical results. Amyloid beta-peptide (25–35) increased caspase-3 expression in the hippocampus, while *Scutellaria baicalensis* stem-leaf total flavonoid decreased caspase-3 expression \( (P < 0.01) \), especially after 100 mg/kg treatment \( (P < 0.01; \text{Figure 8, Table 3}) \).

![Figure 8](image)

**Figure 8** Caspase-3 expression in the hippocampus of amyloid beta-peptide (25–35)-treated rats, detected by western blot analysis. 

SSTF: *Scutellaria baicalensis* stem-leaf total flavonoid.

### Table 3  Effect of *Scutellaria baicalensis* stem-leaf total flavonoid (SSTF) on caspase-3 expression in the hippocampal CA1 region of amyloid beta-peptide (25–35)-treated rats

| Group          | Immunohistochemical staining (absorbance) | Western blot assay (absorbance ratio of caspase-3 to \( \beta\)-actin) |
|----------------|------------------------------------------|-----------------------------------------------------------------------|
| Control        | 22.47±5.14                               | 0.03±0.01                                                             |
| Model          | 126.28±30.58a                            | 0.17±0.03a                                                            |
| Low-dose SSTF  | 60.38±15.49b                             | 0.09±0.02b                                                            |
| High-dose SSTF | 34.78±7.29c                             | 0.07±0.02c                                                            |

Data are expressed as mean ± SD. There were two rats in each group, and four slices (\( \times 400 \)) were selected from each rat for western blot analysis. *P < 0.01, vs. control group; °P < 0.01, vs. model group (one-way analysis of variance, least significant difference between two groups).

*Scutellaria baicalensis* stem-leaf total flavonoid inhibits cytochrome c expression in the hippocampus of amyloid beta-peptide (25–35)-treated rats

Immunohistochemistry demonstrated that cytochrome c was expressed in the cytoplasm of rat hippocampal neurons and occasionally in the cell nuclei, appearing as brownish yellow staining. The number of cytochrome c-positive cells in the hippocampal CA1 region of rats in the model group was increased compared with the control group, and cells were deeply stained. Compared with the model group, cytochrome c-positive cells decreased in number and were lightly stained after low- and high-dose *Scutellaria baicalensis* stem-leaf total flavonoid treatment (Figure 9).

Western blot analysis results were consistent with the immunohistochemical results. Amyloid beta-peptide (25–35) treatment produced an increase in cytochrome c expression in the hippocampus, while *Scutellaria baicalensis* stem-leaf total flavonoid decreased cytochrome c expression \( (P < 0.01) \), especially after 100 mg/kg treatment \( (P < 0.01; \text{Figure 10, Table 4}) \).

### DISCUSSION

Alzheimer’s disease is a neurodegenerative disease characterized by an imbalance in pro-apoptotic and anti-apoptotic processes. *Scutellaria baicalensis* stem-leaf total flavonoid may attenuate learning and memory impairment and cell damage induced by amyloid beta. This function of the traditional Chinese medicine is closely related to its antioxidative effects. In this study, we used the TUNEL assay, immunohistochemistry and
western blot analysis to assess the hippocampal expression of the apoptosis effector molecule caspase-3, the pro-apoptotic gene Bax, cytochrome c and the anti-apoptotic gene Bcl-2. The aim of this study was to explore the effects of Scutellaria baicalensis stem-leaf total flavonoid on amyloid beta-induced damage and apoptosis.

Results of the TUNEL assay showed that amyloid beta triggered apoptosis in the hippocampus.

Immunohistochemical results revealed that caspase-3, Bax and cytochrome c expression levels in the hippocampus increased, while Bcl-2 expression decreased. Western blot analysis suggested that hippocampal expression of Bax, cytochrome c and caspase-3 in the model group was higher than in the control group, while Bcl-2 expression was lower. Collectively, the results indicate that amyloid beta can promote pro-apoptotic protein expression and inhibit anti-apoptotic protein expression.

Table 4. Effect of Scutellaria baicalensis stem-leaf total flavonoid (SSTF) on cytochrome c expression in the hippocampal CA1 region of amyloid beta-peptide (25–35)-treated rats

| Group             | Immunohistochemical staining (absorbance) | Western blot assay (absorbance ratio of cytochrome c to β-actin) |
|-------------------|-------------------------------------------|---------------------------------------------------------------|
| Control           | 25.13±4.37                                | 0.61±0.12                                                     |
| Model             | 124.79±30.78                              | 1.25±0.21                                                     |
| Low-dose SSTF     | 89.27±20.46                               | 0.83±0.16                                                     |
| High-dose SSTF    | 54.18±16.27                               | 0.78±0.15                                                     |

Data are expressed as mean ± SD. There were two rats in each group, and four slices (×400) were selected from each rat for immunohistochemistry. Six rats in each group were taken for western blot analysis. *P < 0.01, vs. control group; #P < 0.01, vs. model group (one-way analysis of variance, least significant difference between two groups).

How does amyloid beta induce neuronal apoptosis? The mitochondrial apoptotic pathway involves various proteins, especially p53, Bax, Bcl-2 and caspase[2, 12]. Caspase-3 is the key factor mediating apoptosis, and its activation contributes to cytochrome c release from mitochondria, mitochondrial damage and neuronal apoptosis[13]. Bcl-2 and Bax are a pair of antagonistic proteins in the Bcl-2 gene family. Bcl-2 has significant anti-apoptotic effects, while Bax promotes apoptosis[14–15]. Bax and Bcl-2 can control the release of cytochrome c by regulating mitochondrial outer membrane permeability[16].

When Bax predominates, the cytochrome c released from the mitochondria may trigger a series of cascade reactions leading to apoptosis. The mitochondrial apoptotic pathway can be divided into caspase-dependent and independent pathways. Mitochondrial amyloid beta can increase the permeability of the mitochondrial outer membrane and promote cytochrome c release, this is called the first phase of release. The released cytochrome c stimulates inositol 1,4,5-triphosphate receptors on the endoplasmic reticulum and promotes unrestricted release of calcium. The intracellular calcium increase leads to a further increase in cytochrome c release, inducing apoptosis, which is independent of caspase[17–18]. Amyloid beta also
decreases complex III (ubiquinone-cytochrome c-reductase) and IV (cytochrome c-oxidase) activity in the mitochondrial respiratory chain[19]. The reduced activity leads to a decrease in ATP, and the excess electrons directly trigger the production of superoxide anion (O$_2^-$) and other reactive oxygen species[20]. Cardiolipin is the only phospholipid in the mitochondrial inner membrane and is the target of reactive oxygen species. Reactive oxygen species may cause myocardial lipid peroxidation, and oxidized cardiolipin has a low binding affinity for cytochrome c[21], thereby leading to complete cytochrome c release, this is called the second phase of release. In the presence of ATP and dATP, cytochrome c mediates the activation of apoptosis protease-activating factor 1, inducing apoptosis complex formation and stimulating caspase-9 activation, which ultimately triggers catalytic maturation and apoptosis[22]. This apoptotic process is dependent on caspase. In this study, Bax expression increased, Bcl-2 expression decreased and cytochrome c was increased in the model group. This evidence suggests that amyloid beta may affect the expression of Bax and Bcl-2, further influencing mitochondrial outer membrane permeability, thereby regulating the release of cytochrome c and inducing neuronal apoptosis that is independent of caspase. In addition, caspase-3 expression was shown to increase. Amyloid beta causes an increase in reactive oxygen species[23]. Therefore, amyloid beta may also trigger cytochrome c release via reactive oxygen species, activating caspase and inducing caspase-dependent apoptosis.

TUNEL, immunohistochemistry and western blot assay results showed that cellular apoptosis decreased significantly, Bax expression decreased, Bcl-2 expression increased, and cytochrome c expression decreased in rats treated with Scutellaria baicalensis stem-leaf total flavonoid. This evidence suggests that Scutellaria baicalensis stem-leaf total flavonoid antagonizes amyloid beta-induced changes to apoptosis-related proteins, and alleviates caspase-independent neuronal apoptosis. Scutellaria baicalensis stem-leaf total flavonoid down-regulates caspase-3 expression and has an antioxidant effect. Thus, the anti-apoptotic effects of Scutellaria baicalensis stem-leaf total flavonoid may be mediated by its antioxidant effect, decreasing reactive oxygen species production and alleviating caspase-dependent apoptosis. Zhu et al[24] found that Hibifolin could inhibit the activation of caspase-3 and caspase-7, decrease Ca$^{2+}$ flow, and reduce DNA damage in cortical neurons induced by amyloid beta. Zhang et al[11] also demonstrated that flavonoid may regulate the expression of Bcl family proteins and have an anti-apoptotic role in the mitochondrial death pathway.

In addition to antioxidation, anti-apoptosis and alleviating neurotoxicity, flavonoid also affects amyloid beta generation and fibrillation to protect neurons. Jung et al[25] found that total flavonoid extracted from Sophora plants can inhibit β-secretase activity. Thus, because amyloid precursor protein produces amyloid beta through β and γ secretase cleavage, inhibiting β-secretase activity can reduce amyloid beta production. Naiki et al[26] studied five kinds of flavonoid in vitro and found that flavonoid inhibits the formation of amyloid beta fibrils and promotes fibrillar instability. Soluble amyloid beta has no toxicity, but it will acquire toxicity after fibrillation. Lemkul et al[27] observed that morin binds to the end of fibrils and functions to block peptide adherence. It penetrates into the hydrophobic center and cleaves at aspartic acid and lysine residues, interfering with backbone hydrogen bonds, ultimately inhibiting fibrillation.

In summary, Scutellaria baicalensis stem-leaf total flavonoid can attenuate neuronal apoptosis induced by amyloid beta. It accomplishes this by modulating the expression of apoptosis-related genes and by having an antioxidant effect. Our findings provide an experimental basis for the application of Scutellaria baicalensis stem-leaf total flavonoid in the prevention and treatment of neurodegenerative diseases.

**MATERIALS AND METHODS**

**Design**
A randomized, controlled, animal experiment.

**Time and setting**
The experiment was performed in the Animal Laboratory of Chengde Medical College, China from June 2009 to June 2010.

**Materials**

**Animals**
Forty healthy, clean, male, Wistar rats, aged 10–12 weeks, weighing 280 ± 20 g, were provided by the Experimental Animal Center of Hebei Province (license No. SYXK (Ji) 2009-0022). All experiments were performed in the specific pathogen free level at 22 ± 2°C and 50 ± 5% humidity, allowing free access to water and food. All procedures were in accordance with the Guidance Suggestions for the Care and Use of...
**Laboratory Animals**, formulated by the Ministry of Science and Technology of China\(^{[28]}\).

**Drugs**

*Scutellaria baicalensis* stem-leaf total flavonoid was provided by the Institute of Traditional Chinese Medicine, Chengde Medical College, China. In brief, stems and leaves were crushed into pieces and extracted two or three times in aqueous solution. The supernatant was acidized, isolated, adsorbed, desorbed, condensed and dried to prepare a brown powder of 61.88% purity\(^{[3]}\).

*Scutellaria baicalensis* stem-leaf total flavonoid was dissolved in distilled water and adjusted for pH 7.0.

Amyloid beta-peptide (25–35) (purity 97%; Sigma, St. Louis, MO, USA) was tested with high performance liquid chromatography. Amyloid beta-peptide (25–35), 1 mg, was dissolved in 500 μL sterile saline to prepare a 2 g/L solution, which was stored at −20°C. Prior to the experiment, the solution was incubated at 37°C for 5–7 days.

**Methods**

**Drug intervention**

Rats in the control and model groups were intragastrically given distilled water, and those in the low-dose and high-dose treatment groups were treated with *Scutellaria baicalensis* stem-leaf total flavonoid, 50 or 100 mg/kg per day, respectively, for 20 consecutive days. On day 8 after administration, 5 μL amyloid beta-peptide (25–35), 10 μg, was injected into the bilateral hippocampus of rats in the control group, while 5 μL amyloid beta-peptide (25–35), 10 μg, was injected into the bilateral hippocampal CA1 region of rats (anteroposterior: −3.5 mm; mediolateral ± 2 mm; dorsoventral: 2.7 mm\(^{[25]}\)) in the three remaining groups.

**Tissue preparation**

All animals were killed after 20 days of intragastric administration, and four rats from each group were selected for preparation of brain tissue paraffin specimens. Blood samples were collected under anesthesia, and 250 mL saline was perfused via the left ventricle. Brain tissue was fixed in PB solution, 250 mL, containing paraformaldehyde, for 1 hour. Then, the bilateral hippocampi were cut along the coronal plane, and the tissue block was trimmed and fixed in 4% paraformaldehyde overnight. Brain tissue was rinsed with PBS, dehydrated in graded ethanol, cleared with xylene, and embedded in paraffin. Two pieces of tissue were embedded in a piece of wax and cut into serial coronal slices of 5 μm thickness, containing the hippocampus, for TUNEL and immunohistochemical staining.

**TUNEL staining for hippocampal cell apoptosis**

The paraffin slices were dewaxed, hydrated and assayed according to instructions in the apoptosis detection kit (Roche, Basel, Switzerland). Slices were incubated with reaction liquid and converter-POD, developed with 3,3’-diaminobenzidine, counterstained with hematoxylin, dehydrated in graded ethanol, cleared with xylene, and mounted with neutral gum. Brain tissue was observed under an optical microscope (Olympus, Tokyo, Japan). A cell with its cytoplasm or nucleus stained brown-yellow indicated an apoptotic cell.

**Immunohistochemical staining for Bax, Bcl-2, caspase-3 and cytochrome c expression in the rat hippocampus**

Brain slices were dewaxed with xylene, hydrated with gradient ethanol, and placed in a 95°C water bath for antigen retrieval. Slices were incubated with rabbit anti-Bax polyclonal antibody (1:100), rabbit anti-caspase-3 polyclonal antibody (1:200), rabbit anti-Bcl-2 polyclonal antibody (1:75) or mouse anti-rat cytochrome c monoclonal antibody (1:75) at 37°C for 2.5 hours. All antibodies were purchased from Santa Cruz Biotechnology, Santa Cruz, CA, USA. Then, slices were incubated with horseradish peroxidase-conjugated goat anti-rabbit/ mouse IgG (1:3 000; Beijing Zhong Shan-Golden Bridge Biological Technology Co., Ltd., Beijing, China) at 37°C for 15 minutes, developed with 3,3’-diaminobenzidine, counterstained with hematoxylin, and mounted with neutral gum. Brain tissue was observed under the light microscope. Negative control was incubated with PBS, rather than antibodies.

**Image analysis**

Two rats in each group were selected for preparation of brain tissue paraffin sections for TUNEL and immunohistochemical staining. Four consecutive sections from the hippocampal CA1 region were observed by light microscopy, and the absorbance was measured using Image-Pro Plus software (Media Cybernetics, Inc., Bethesda, MD, USA) under 400× magnification and averaged.

**Western blot analysis for Bax, Bcl-2, caspase-3 and cytochrome c expression in the rat hippocampus**

Hippocampal tissue, 0.15–0.20 g, from six rats in each group was incubated with 1 mL PBS-NaF, ground on ice, centrifuged, and homogenized with RIPA lysate (Beyotime Institute of Biotechnology, Shanghai, China). Protein content was measured according to instructions...
in the bicinechonic acid kit (BestBio, Shanghai, China). Samples were loaded onto sodium dodecyl sulfate polyacrylamide gel, electrophoresed, transferred onto a polyvinylidene difluoride membrane (IPVH00010; thickness: 0.45 μm, 0.22 μm; Millipore, Billerica, MA, USA), and blocked overnight. Then, slices were incubated with anti-Bax polyclonal antibody (1:200), rabbit anti-caspase-3 polyclonal antibody (1:200), rabbit anti-Bcl-2 polyclonal antibody (1:100), mouse anti-rat cytochrome c monoclonal antibody (1:200) and mouse-rat β-actin monoclonal antibody (1:1 000; Santa Cruz Biotechnology) at room temperature for 2 hours. Afterwards, slices were incubated with horseradish peroxidase-conjugated goat anti-rabbit IgG (1:3 000) and goat anti-mouse IgG (1:5 000) for 1 hour. Images were developed and fixed using enhanced chemiluminescence (Applygen Technologies Inc., Beijing, China). Absorbance was measured with Gelpro4 software, and β-actin was taken as the internal reference.

**Statistical analysis**

SPSS 11.5 software (SPSS, Chicago, IL, USA) was used for one-way analysis of variance, and two groups were compared using the least significant difference test method. Measurement data were expressed as mean ± SD. A P value < 0.05 was regarded to indicate a significant difference.

**Acknowledgments:** We are grateful to Professor Jianxin Zhang from the Hebei Academy of Medical Sciences; Professor Jiming Tong from the Institute of Traditional Chinese Medicine of Chengde Medical College; Xin Li, Yong Yan, Xiaoguang Wu, Long Chen, Qian Xu and Xiaochun Zhou from the Institute of Basic Sciences of Chengde Medical College; Professor Shumin Zhao and all staff from the Department of Electron Microscopy of Chengde Medical College, China; Liquan Ren, Lixin Mei and Yingchun Zhang from the Department of Pathology of Chengde Medical College, China; Professor Lixin Sun from the Central Laboratory of Affiliated Hospital of Chengde Medical College, China; and Xiaoping Jin from the Department of Pathology in the Affiliated Hospital of Chengde Medical College, China for help with experimental guidance.

**Funding:** This study was supported by grants from Hebei Provincial Science and Technology Bureau, No. 08276101D-21.

**Author contributions:** Ruiting Wang was responsible for the study design and implementation. Xingbin Shen performed morphological experiments. Enhong Xing performed molecular biology experiments. Lihua Guan participated in animal experiments and immunohistochemistry. Lisheng Xin conducted statistical processing. Ruiting Wang wrote the manuscript. All authors approved the final version of the manuscript.

**Conflicts of interest:** None declared.

**Ethical approval:** This study was approved by the Animal Ethics Committee of Chengde Medical College in China.

**Author statement:** The manuscript is original, has not been submitted to or is not under consideration by another publication, has not been previously published in any language or any form, including electronic, and contains no disclosure of confidential information or authorship/patent application/funding source discriptions.

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(Reviewed by Patel B, Stow A, Liu P, Ma XL) (Edited by Wang LM, Yang Y, Li CH, Song LP)