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Ameliorative effects and possible molecular mechanisms of action of fibrauretime from *Fibraurea recisa* Pierre on D-galactose/ AlCl_3 -mediated Alzheimer's disease

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Fibrauretime is one of the main active ingredients from the rattan stems of *Fibraurea recisa* Pierre. It exhibits a series of significant pharmacological effects. The present study aimed to evaluate the potential anti-Alzheimer's disease (AD) effects of fibrauretime on a D-galactose/ AlCl_3 -induced mouse model, and the underlying mechanisms of action were further investigated for the first time. Our results showed that pretreatment with fibrauretime significantly improved the ability of spatial short-term working memory in the model mice during the Y-maze test, as well as the abilities of spatial learning and memory during the Morris water maze. The levels of brain tissue amyloid ($\text{A}\beta$), P-Tau, Tau and acetylcholinesterase (AChE) were evidently increased in D-galactose/ AlCl_3 -intoxicated mice, and these effects were reversed by fibrauretime. In contrast, a significant increase in the levels of the neurotransmitter acetylcholine (ACh) and choline acetyl transferase (ChAT) was observed in the fibrauretime-treated groups compared with the model group. Neuronal oxidative stress, evidenced by increased malondialdehyde (MDA) and nitric oxide (NO) levels and a decline in glutathione (GSH), catalase (CAT) and superoxide dismutase (SOD) activity, was significantly alleviated by fibrauretime pretreatment. The suppression of the neuroinflammatory response by fibrauretime was realized not only by the decrease in the levels of tumour necrosis factor- α (TNF- α) and interleukin-1 β (IL-1 β) in the brain tissues and by the enzyme-linked immunosorbent assay (ELISA) but also by the protein expression levels of nuclear factor- κB (NF- κB), cyclooxygenase-2 (COX-2), and inducible nitric oxide synthase (iNOS), which were measured by immunohistochemistry and western blotting. In addition, the protein expression levels of inflammatory factors interleukin-33 (IL-33) and ST2 in the brain tissues were detected by immunohistochemistry. Furthermore, the effects of western blotting demonstrated that the administration of fibrauretime significantly suppressed the protein expression levels of caspase-3, cleaved caspase-3, and Bax and increased the protein expression levels of Bcl-2, and the results of the H&E and TUNEL assay all suggested the inhibition of apoptosis in the neurons. The results clearly suggest that the underlying molecular mechanisms of action of the fibrauretime-mediated alleviation of D-galactose/ AlCl_3 -induced Alzheimer's disease may involve antioxidant, anti-inflammatory, and anti-apoptotic effects.

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1 Introduction

Alzheimer's disease (AD) is the most common type of dementia among the elderly, and AD has been prevalent with the rapidly growing global elderly population.¹ AD is a fatal neurodegenerative disease clinically characterized by progressive decline in cognitive functions such as memory loss and behavioural disturbances.² The neuropathologic

hallmarks of AD include the presence of extracellular $\text{A}\beta$ amyloid plaque (AP) deposition and the intracellular formation of neurofibrillary tangles (NFTs) in the brain, together with synaptic dysfunction characterized neurochemically by a consistent deficit in cholinergic neurotransmission and a significant loss of neuronal cells.³ These lesions contain hyperphosphorylated tau and an excess of reactive glia in the AD brain.^{4,5} The characteristics of the ageing model for D-galactose-induced AD mice is close to the pathological changes of clinical AD patients. A large amount of aluminium ions may induce AP deposition and formation of NFTs and is a common damage model of the cholinergic system in AD. AD

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model mice may be effectively induced by D-galactose combined with AlCl_3 (D-galactose/ AlCl_3).^{6–8}

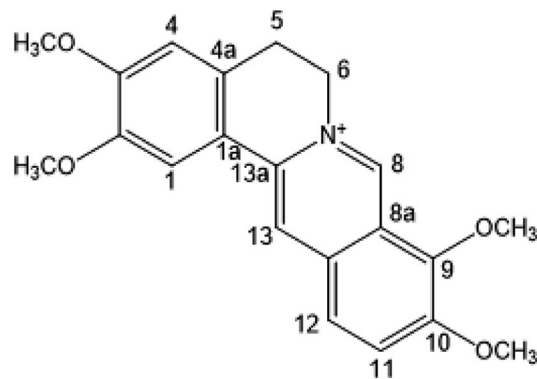
The mechanisms underlying the progression of AD remain complicated. The continuous generation and abnormal accumulation of A β in the brain causes microglial dysfunction by reducing A β receptors, increasing the production of inflammatory mediators,⁹ including TNF- α , IL-1 β , NF- κ B, COX-2 and iNOS. Therefore, chronic neuroinflammation induced by microglia contributes to pathological progression and symptom severity, usually in the late stages of AD. Interleukin-33 (IL-33) is a new member of the IL-1 family, whose specific receptor is ST2.¹⁰ IL-33 is constitutively expressed in the central nervous system (CNS), may induce inflammatory molecule release from glial cells and plays an important role in the pathogenesis of AD.¹¹

Oxidative stress is an imbalance between the oxidation and reduction reactions, resulting in the production of reactive oxygen species (ROS).¹² Oxidative stress plays a very important role in the etiopathogenesis of AD.¹³ Free radicals can be produced by microglial activation and generated by β -amyloid plaques. Additionally, the activation of a series of signalling protein pathways such as inflammatory pathway nuclear factor- κ B (NF- κ B)¹⁴ can be triggered by ROS-mediated oxidative stress. Neuronal apoptosis is believed to lead to AD, and apoptosis also plays a vital role in the pathophysiology of AD.^{15,16} Apoptosis is a sophisticated biological mechanism involving a wide variety of anti-apoptosis proteins such as Bcl-2 and pro-apoptotic protein such as Bax.¹⁷ Caspase-3, the effector caspase and final executor of apoptosis, can cleave cytoskeletal and nuclear proteins.¹⁸

Several studies have focused currently on natural ingredients from traditional herbal medicines to evaluate novel therapeutic drugs for the therapy of AD. Various natural ingredients, including luteolin, tetrahydropalmatine and berberine have been shown ameliorative effects on AD *in vivo* experiments.^{19–21}

The rattan stems of *Fibraurea recisa* Pierre is rich in resources and mainly distributed in Yunnan and Guangxi, China, known as natural antibiotics, have traditionally been used in the treatment of gynaecological inflammation and surgical infection. The rattan stems contains a kind of isoquinoline alkaloid, one of the main active ingredients, fibrauretine. More significant pharmacological activities of fibrauretine have been discussed in the previous study, involving antitumour,^{22,23} anti-osteoarthritis,²⁴ antiviral,²⁵ neuroprotective,^{26,27} antidepressant²⁸ effects and lowering blood lipids.²⁹

In this paper, fibrauretine was extracted and purified from the rattan stems of *Fibraurea recisa* Pierre with 60.75% extraction rate. The anti-AD effects of fibrauretine against D-galactose/ AlCl_3 -induced AD surveyed on a mouse model were proposed for the first time by us. The new discovery has been authorized by the State Intellectual Property Office of China (ZL201510253034.6). In addition, the potential molecular mechanisms underlying these anti-AD effects are discussed, involving antioxidant, anti-inflammatory, and anti-apoptotic activity.



fibrauretine

2 Materials and methods

2.1 Preparation of fibrauretine

The rattan stems of *Fibraurea recisa* Pierre were purchased in September 2015, in Kunming of Yunnan Province, China and were identified by Prof. Lianxue Zhang. The voucher specimen was deposited at the laboratory.

The air-dried rattan stems (2 kg, 60 mesh) were extracted two times with 70% ethanol by reflux extraction, each time for 2 hours. The extracts were filtered, combined and concentrated under reduced pressure to give 228.3 g of extract. The extracts were then dissolved in water and passed through a pre-treated D101 macroporous resin column ($\Phi 1000 \times 100$ mm), washed with 3 column volumes of water, and then eluted with 8 column volumes of 40% ethanol. The ethanol eluate was collected, the ethanol was recovered, and the eluate was then dried to yield total alkaloids (50.6 g). Total alkaloids (10 g) were dissolved in 95% ethanol, chromatographed on a basic alumina column (200 g, $\Phi 880 \times 100$ mm) and eluted with 95% ethanol for elution at atmospheric pressure. The alkaloids were detected by thin layer chromatography (TLC) and separated to give compound A and component I. Component I was subjected to gradient elution with methanol–water (10 : 90–50 : 50) on an ODS column ($\Phi 420 \times 36$ mm) to afford fibrauretine (24.3 g). The purity of fibrauretine was determined by HPLC (purity > 98%), and the structure was identified by ¹H-NMR and ¹³C-NMR.³⁰

2.2 Animals and experimental protocol

ICR mice (male, weighing 18–22 g, SPF grade) were provided by YISI Experimental Animals Co., Ltd. (Certificate no. SCXK-2016-0003, Jilin, China). Animals were kept under standard laboratory conditions: room temperature at 23 ± 2 °C, humidity of $60 \pm 10\%$. The mice were allowed free access to food and water during the experiments. All experimental procedures were approved by the Institutional Animal Ethical Committee and were authorized by the Animal Care Committee of Jilin Agricultural University (2016-01-Permit number: ECLA-JLAU 2016-016), the reference is Guide for the Care and Use of Laboratory Animals (Eighth Edition, the National Academies Press Washington, D.C.) and AVMA Guidelines for the Euthanasia of



Animals: 2013 Edition (American Veterinary Medical Association).

After acclimation for 3 days, 10 mice were randomly selected as a control group. The remaining 80 mice received intraperitoneal injections with D-galactose at a dose of 120 mg kg⁻¹ and were administered AlCl₃ intragastrically at a dose of 20 mg kg⁻¹ daily for 40 continuous days to induce AD in mice. The control group mice were injected intraperitoneally and administered intragastrically with the same amount of saline. From the 35th to the 40th day, after 1 h of being given D-galactose/AlCl₃, the memory behaviour of the mice was trained using the Y-type maze test for five continuous days. The eligible mice were randomly divided into 5 groups (*n* = 10) and were then treated for the second forty days according to methods mentioned above. All mice were administered intragastrically with different

drugs from the 41st day for 40 continuous days. According to our previous study of the LD₅₀ of fibrauretime (4.8 g kg⁻¹) and the preliminary experiment, the groups for fibrauretime administration were, respectively, given different dosages (80, 160, 320 mg kg⁻¹). The positive control group was treated with donepezil hydrochloride (1 mg kg⁻¹). Fibrauretime and donepezil hydrochloride were, respectively, suspended in 0.05% carboxymethylcellulose sodium (CMC-Na) before being used. The control group mice and the model group mice received intragastrical injections with 0.05% CMC-Na. The administration operators were blind to the tested materials, and the testers were also blind to the drug treatment.

All mice were weighed every three days. Eighty days later, after 1 h from the last administration, each group of mice was subjected to the Morris water maze and the Y-maze test and

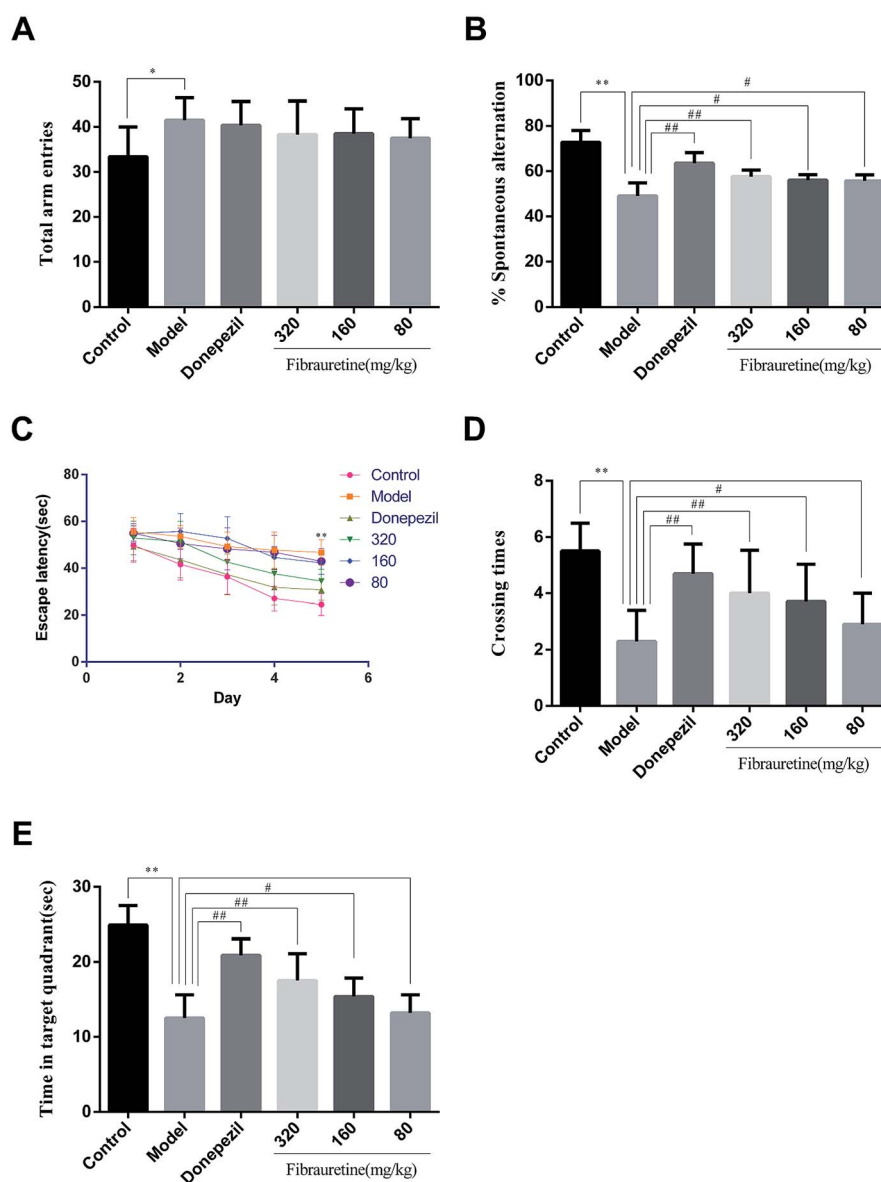


Fig. 1 Effect of fibrauretime and donepezil on total arm entries (A); spontaneous alternation (B); escape latency (C); crossing times (D); time in target quadrant (E) from D-galactose/AlCl₃-induced AD mice. All data were expressed as the mean \pm SD, *n* = 10. **p* < 0.05, ***p* < 0.01 vs. control group; #*p* < 0.05, ##*p* < 0.01 vs. model group.



then was sacrificed under general anaesthesia. Blood and sample collection was performed immediately. The brain tissue was promptly frozen in liquid nitrogen and stored at $-80\text{ }^{\circ}\text{C}$ until analysis, and the brain tissue was fixed in 10% neutral buffered formalin. Subsequently, serum samples were separated using a refrigerated centrifuge at $3000 \times g$ and stored at $-20\text{ }^{\circ}\text{C}$ until used for subsequent analysis.

2.3 Y-maze test

The mice were evaluated for short-term spatial memory by a Y-maze test as previously described.³¹ The Y-maze consisted of three identical arms (60 cm in length, 10 cm in width, and 15 cm in height), 120 degrees to each other. The mice were individually placed at the end of either arm and allowed to move freely through the maze within 5 minutes. Mice actively explore the maze, enter each arm in order, and remember which arm they have already entered. Spontaneous alternation is defined as the continuous entry of three arms (*e.g.*, 1/2/3 or 1/3/2), and

the spontaneous alternation rate is the ratio of the actual number of alternations to the number of possible alternations (the total number of arms minus 2) multiplied by 100%.

2.4 Morris water maze

The spatial learning and memory of mice was tested by a water maze as described in.³² The Morris water maze consisted of a 100 cm diameter pool that was filled with water ($22\text{ }^{\circ}\text{C}$). The pool was divided into four equal quadrants, and there were visual cues in each quadrant. Mice were accepted to the hidden platform test from day 1 to day 4, with a probe trial on the fifth day. In the hidden platform test, the platform was placed 1 cm underwater, and each time, the mice were randomly placed in different starting positions; they were given 60 seconds to climb the hidden platform. If no platform could be found within 60 seconds, mice were guided to the platform and allowed to remain on the platform for 10 seconds. In the probe trial, the platform was removed, and mice were put into the pool to swim

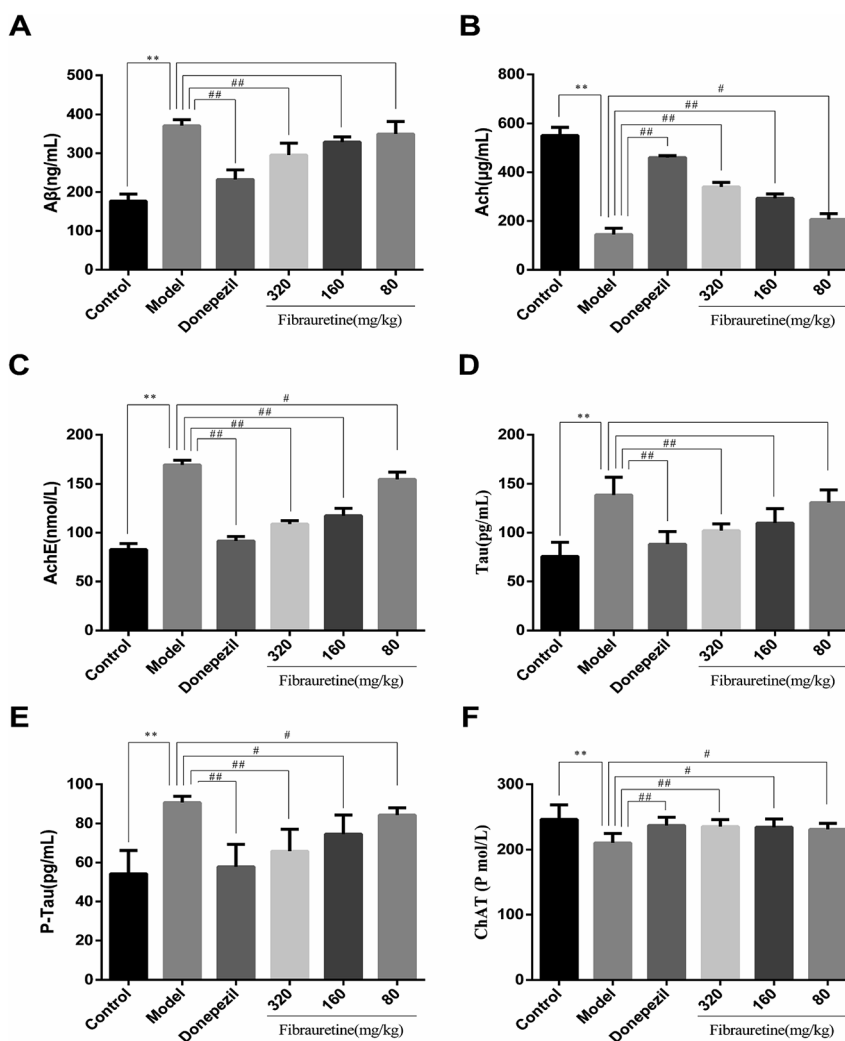


Fig. 2 Effects of fibrauretime and donepezil on the level of brain A β (A); the level of brain Ach (B); the level of brain AchE (C); the level of brain Tau (D); the level of brain P-Tau (E); and the level of brain ChAT (F) in D-galactose/AICl₃-induced AD. All data are expressed as the mean \pm SD, $n = 10$. * $p < 0.05$, ** $p < 0.01$ vs. control group; # $p < 0.05$, ## $p < 0.01$ vs. model group.



for 60 seconds. The memory consolidation 24 hours after the last hidden-platform trial was assessed. The time the mouse spent in the target quadrant and the number of crossings were detected.

2.5 Assessment of biochemical parameters

The brain tissue levels of A β , AchE, Ach, Tau, P-Tau, ChAT, GSH, TNF- α and IL-1 β , as well as the serum levels of MDA, NO, CAT and SOD, were measured by ELISA. Brain tissue from each group was homogenized on ice, serum samples were segregated using a refrigerated centrifuge at 3000 \times g, then all samples were tested immediately by the ELISA kit (all purchased from R&D systems) according to the manufacturer's instructions.

2.6 H&E staining assay

Histopathological examination was performed as previously described.³³ Briefly, brain tissues were fixed in 4%

formaldehyde solution. After deparaffinization with xylene, the tissues were dehydrated with different concentrations of ethanol and cut into 5 μ m-thick slices. Then, all slices were stained with H&E staining solution and PAS reagents for histopathological analysis. The results were detected by optical microscope (Olympus, BX51T-PHD-J11, Japan).

2.7 TUNEL staining assay

The TUNEL staining assay was conducted to measure the extent of apoptosis in the brain after D-galactose/AlCl₃ exposure. TUNEL evaluation was typically carried out as follows.³⁴ First, the brain sections (5 μ m-thick) were incubated with 100 μ L of 20 μ g mL⁻¹ proteinase K solution at room temperature for 15 min. Next, the sections were incubated with 100 μ L of 3% H₂O₂ for 10 min and washed with PBS two times. Then, the TUNEL reaction mixed solution was added to the sections, and the sections were put into a humid chamber at 37 $^{\circ}$ C for 1 h.

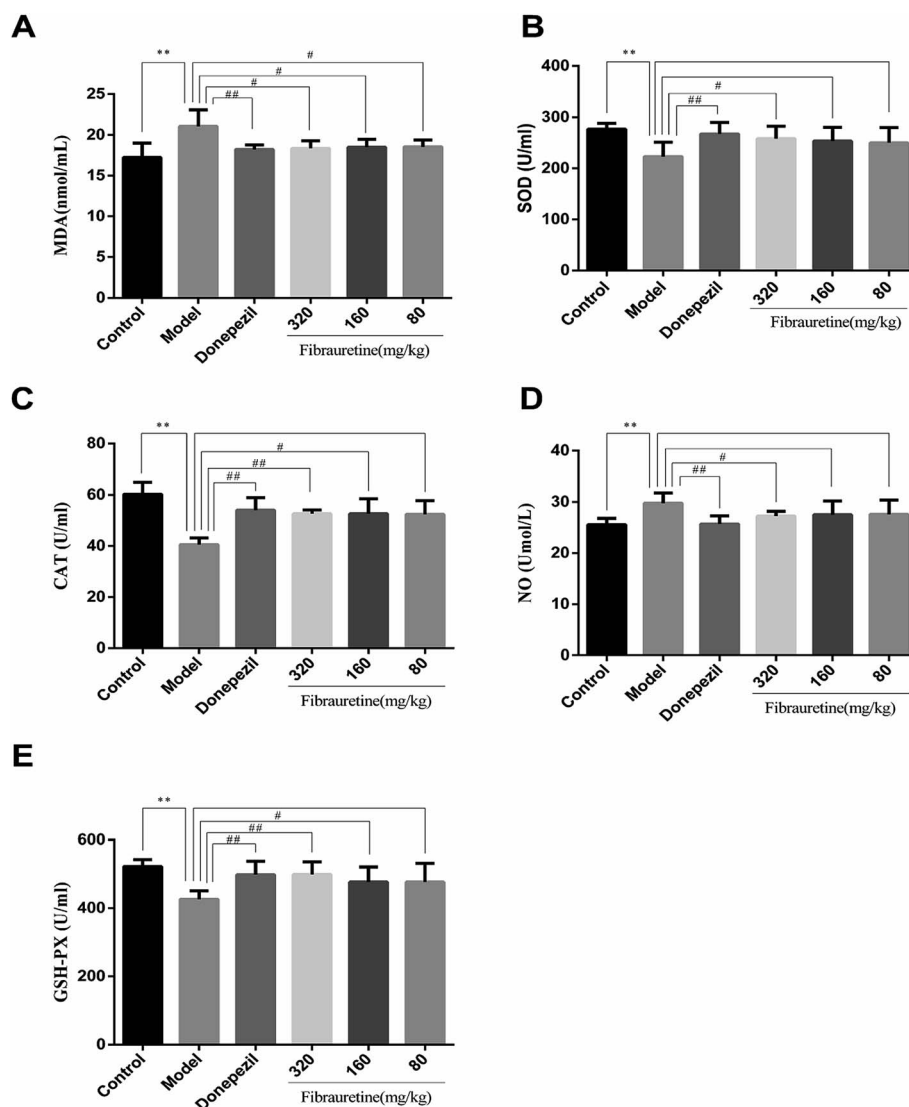


Fig. 3 Effects of fibrauretime and donepezil on the levels of malondialdehyde (MDA) (A); peroxide dismutase (SOD) (B); catalase (CAT) (C); nitric oxide (NO) (D); and glutathione (GSH) (E) in D-galactose/AlCl₃ induced AD. All data are expressed as the mean \pm SD, $n = 10$. * $p < 0.05$, ** $p < 0.01$ vs. control group; # $p < 0.05$, ## $p < 0.01$ vs. model group.



Subsequently, the sections were incubated with 50 μ L POD for 30 min after washing three times with PBS. Finally, the slices were counterstained with haematoxylin. TUNEL-positive cells were visualized with an Olympus microscope (Olympus BX51, Japan).

2.8 Immunohistochemistry (IHC) analysis

IHC analysis was performed as mentioned earlier.³⁵ First, 5 μ m-thick brain sections were de-paraffinized and hydrated with aqueous alcohol solution. The sections were blocked with endogenous peroxidases for 10 min at room temperature, followed by rinsing 3 times in PBS (0.01 M, pH 7.4) for 3 min. Then, the sections were put into citrate buffer solution (0.01 M, pH 6.0) in a microwave oven (medium high temperature) for 15 min. When the temperature came to 25 $^{\circ}$ C, the sections were washed three times with PBS. Next, the slides were incubated with 1% BSA for 1 h and incubated in a humidified chamber at 4 $^{\circ}$ C overnight with initial antibodies involving NF- κ B (1:200), Bax (1:200), Bcl-2 (1:200), iNOS (1:200), COX-2 (1:200), IL-33 (1:200) and ST2 (1:200), followed by HRP-conjugated secondary antibody for 30 min. The sections were incubated with labelled streptavidin-biotin for 30 min. Finally, the sections were followed by DAB staining and counterstained with haematoxylin. The sections were dehydrated with alcohol and cleared with xylene. The immunostaining results were analysed by light microscopy (Olympus BX51, Japan).

2.9 Western blotting analysis

Western blotting analysis was carried out as mentioned previously.³⁶ First, brain tissue samples from each group were homogenized with lysis buffer plus 1 mM PMSF and protease inhibitor. Protein concentrations were determined using a BCA protein assay kit. The proteins (20 μ g per lane) from each sample were separated with 12% SDS-PAGE and transferred to a polyvinylidene fluoride (PVDF) membrane. Then, the

membrane was blocked with 5% fat-free milk in tris-buffered saline (TBS) containing 0.1% Tween-20 for 2 h in a room temperature environment, after being incubated with primary antibodies at 4 $^{\circ}$ C overnight. Caspase-3, cleaved caspase-3, Bax, Bcl-2, NF- κ B, COX-2 and iNOS were probed with the corresponding primary antibodies (1:1000), followed by incubation with HRP-conjugated anti-mouse anti-body or HRP-anti-rabbit antibody. Finally, the blots were inspected by the Emitter Coupled Logic (ECL) substrate with a KODAK Image Station 4000 MM (Carestream Health, Inc., New Haven, CT, USA).

2.10 Statistical analysis

All data were expressed as the means \pm standard deviation (SD) of ten animals in each group and were characterized by a one-way analysis of variance (ANOVA) using SPSS17.0 (SPSS, Chicago, IL, USA). Values of $p < 0.05$ were considered significant, and $p < 0.01$ was considered highly statistically significant.

3 Results

3.1 Effects of fibrauretime on the abilities of learning and memory in D-galactose/AlCl₃-treated mice

The behaviour effect of fibrauretime on D-galactose/AlCl₃-induced AD mice was assessed using a Y-maze test and a Morris water maze test. The results are shown in Fig. 1.

The results from the Y-maze test showed that a single treatment with D-galactose/AlCl₃ raised the total arm entries in the model group compared with the control group ($p < 0.05$), and there was no significant difference between the model group and the fibrauretime-administered group ($p > 0.05$), and there wasn't any significant difference between the model group and the donepezil group in the same way (Fig. 1A). Compared with the control group, the spontaneous alternation rate of mice in the model group was significantly decreased ($p < 0.01$). However, the spontaneous alternation rate was significantly increased by donepezil ($p < 0.01$)

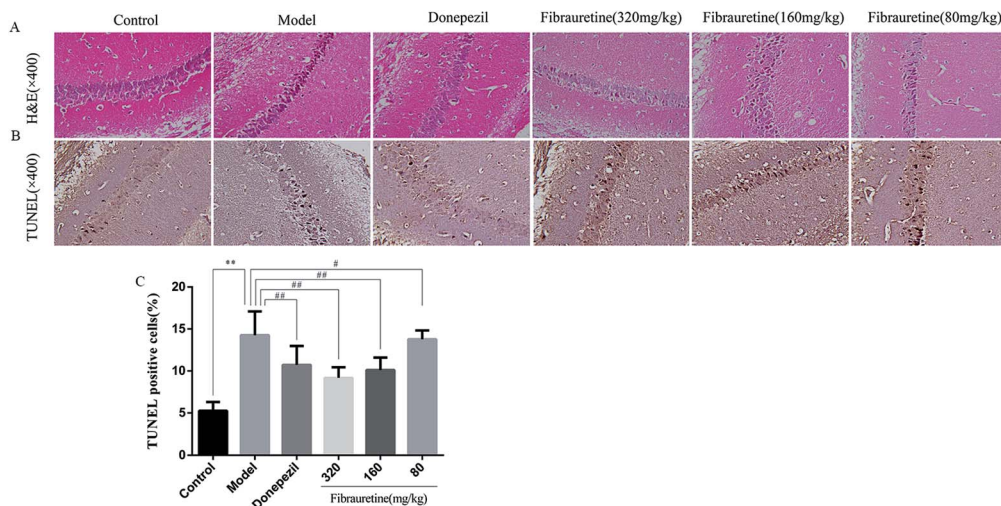


Fig. 4 Morphological changes in D-galactose/AlCl₃ induced AD mice. Brain tissue sections stained by H&E ($\times 400$) (A); brain tissue sections stained by TUNEL ($\times 400$) (B); the presence of TUNEL positive cells measured by the image analyser (C); all data are expressed as the mean \pm SD, $n = 10$. * $p < 0.05$, ** $p < 0.01$ vs. control group; # $p < 0.05$, ## $p < 0.01$ vs. model group.



and fibrauretime at doses of 80, 160 and 320 mg kg⁻¹ in a dose-dependent manner ($p < 0.05$ or $p < 0.01$), as shown in Fig. 1B.

In the water maze experiment, the spatial learning ability of the mice was assessed by the escape latency. The mean escape latencies are shown in Fig. 1C. The results indicate that the escape latency in the model group mice was significantly longer compared with the control group mice ($p < 0.01$). The escape latencies were significantly reduced by donepezil ($p < 0.01$), Similarly, the escape latencies were reduced by fibrauretime at doses of 80, 160 and 320 mg kg⁻¹ ($p < 0.05$ or $p < 0.01$).

Spatial memory retention is assessed by the time spent in the target quadrant and the number of crossings in the target quadrant in the exploratory experiment. Compared with the control group, the time spent in the target quadrant and the number of crossings in the target quadrant in the model group mice were both significantly decreased ($p < 0.01$). In the same way, these changes in the time spent in the target quadrant and the number of passages through the target quadrant were significantly increased by donepezil ($p < 0.01$) and dose-dependently increased by fibrauretime at doses of 80, 160 and 320 mg kg⁻¹ ($p < 0.05$ or $p < 0.01$), as shown in Fig. 1D and E.

3.2 Effects of fibrauretime on biochemical parameters related to AD in mice

The brain levels of A β , AchE, Tau, P-Tau, ChAT and Ach were determined by ELISA assay. The levels of these six biochemical parameters are closely related to AD. They play very important roles in the process of AD. As shown in Fig. 2, compared with the control group, the brain levels of A β , AchE, Tau and P-Tau in the model group were significantly increased ($p < 0.01$), and the brain levels of ChAT and Ach were significantly decreased ($p < 0.01$). A β , AchE, Tau and P-Tau levels of donepezil-administered mice were significantly decreased ($p < 0.01$). And the levels of A β , AchE, Tau and P-Tau in the fibrauretime-administered mice were significantly decreased in a dose-dependent manner ($p < 0.05$ or $p < 0.01$). On the contrary, the levels of ChAT and Ach in the brains of donepezil were significantly increased ($p < 0.01$). And ChAT, Ach levels of fibrauretime mice were significantly increased in a dose-independent manner ($p < 0.05$ or $p < 0.01$). The data above indicated that fibrauretime has an obvious anti-AD effect.

3.3 Effects of fibrauretime on the oxidative stress of the brain in D-galactose/AlCl₃-treated mice

As previously described, oxidative stress injury participated in the mechanisms of AD.¹³ As indicated in Fig. 3, D-galactose/AlCl₃ treatment caused a conspicuous reduction in glutathione (GSH) level, superoxide dismutase (SOD) activity and catalase (CAT) activity, accompanied by an increase in malondialdehyde (MDA) content and nitric oxide (NO) content, compared with the control group ($p < 0.01$). However, compared with the model group, the administration of fibrauretime reduced MDA content, NO content, and recovered antioxidant capacity, as indicated by the increase in the GSH level, SOD activity and CAT activity in a dose-independent manner ($p < 0.05$ or $p < 0.01$). So did the donepezil-treated group at dose of 1 mg kg⁻¹. These data suggested that fibrauretime alleviated oxidative damage in the

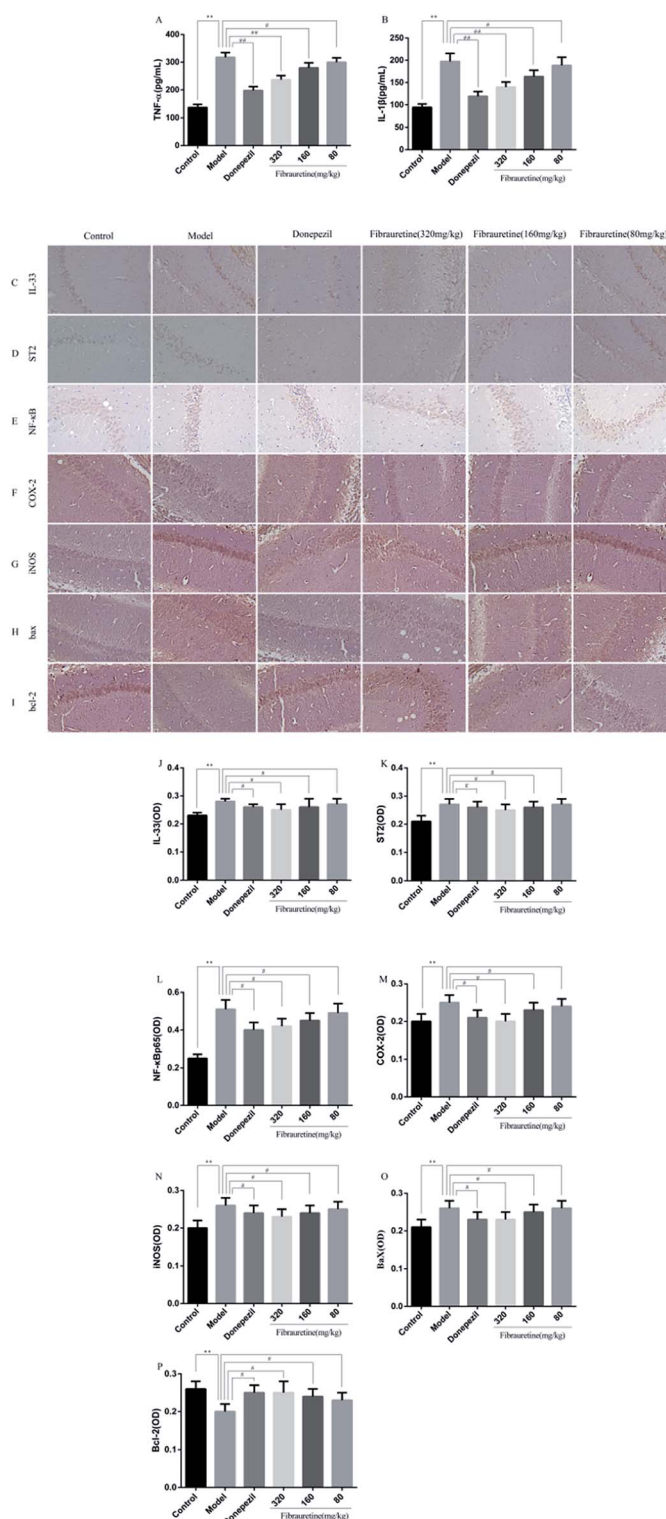


Fig. 5 Effects of fibrauretime and donepezil on the levels of TNF- α (A); IL-1 β (B). Effects of fibrauretime and donepezil on the expression of: IL-33 (C); ST2 (D); NF- κ B (E); COX-2 (F); iNOS (G); Bax (H) and Bcl-2 (I). The column chart shows the antibodies stained area of: IL-33 (J); ST2 (K); NF- κ B (L); COX-2 (M); iNOS (N); Bax (O) and Bcl-2 (P). The protein expression was determined by immunohistochemistry in brain tissues from all groups. * $p < 0.05$, ** $p < 0.01$ vs. control group; # $p < 0.05$, ## $p < 0.01$ vs. model group.



serum and brain tissues by up-regulating antioxidant enzyme activity.

3.4 Effects of fibrauretime on histopathological changes of the brain in D-galactose/ AlCl_3 -treated mice

According to the brain tissue sections in D-galactose/ AlCl_3 -induced AD mice, the hippocampal neuron cell growth and shape was observed by H&E staining (Fig. 4A). The results

showed that there were fewer neurons of the hippocampus in the model group arranged heterocentrically, in which cell body, pyknotic and deeply stained nuclei shrank and the elongated axon could be found. However, these morphological changes in fibrauretime-treated group could be ameliorated. In the hippocampal region of fibrauretime 320 mg kg^{-1} and 160 mg kg^{-1} -treated groups, the neurons were arranged regularly and increased, and the improvement in the structure of the hippocampus neurons was obvious. The same result as the

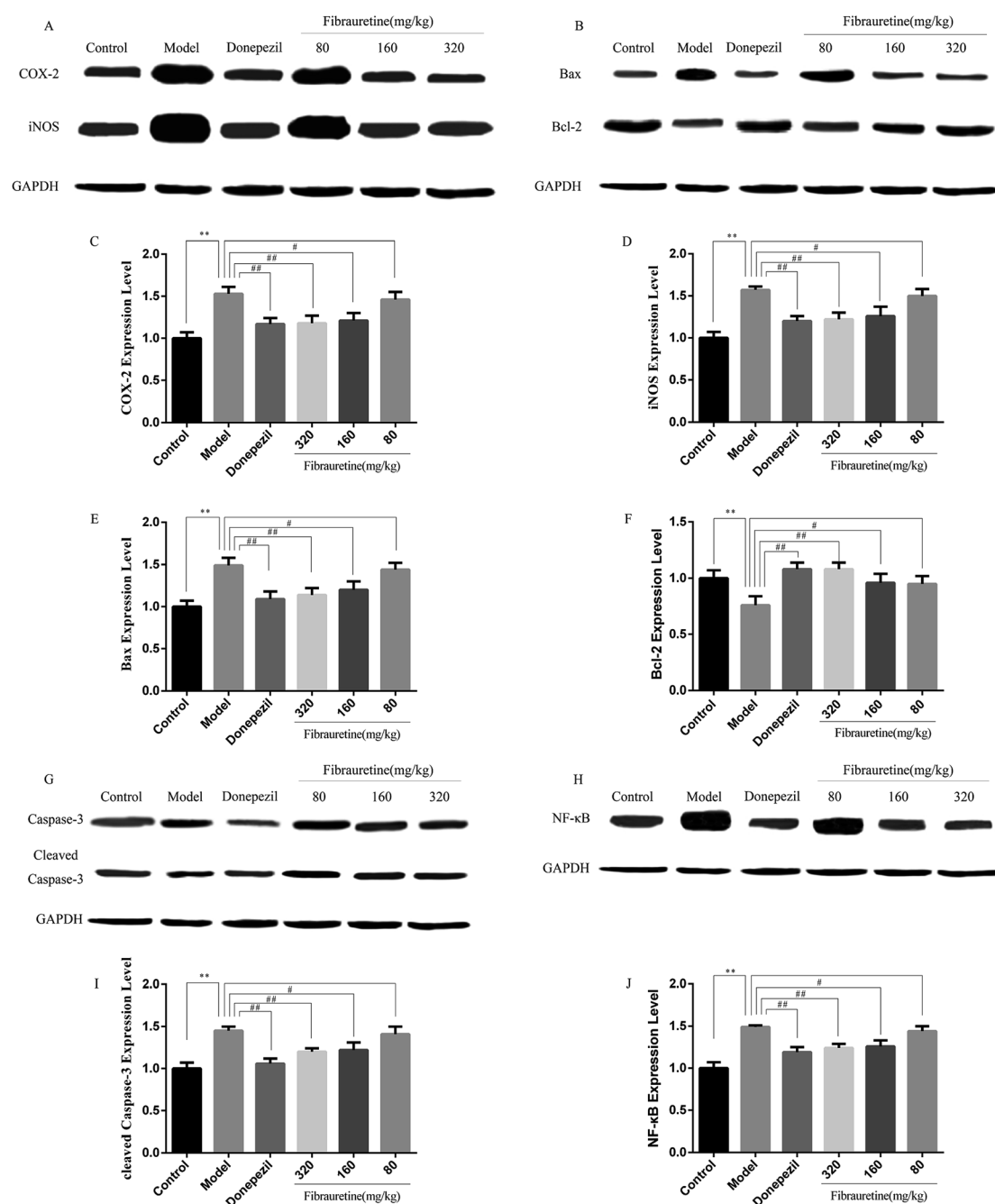


Fig. 6 Effects of fibrauretime and donepezil on the protein expression of COX-2, iNOS (A); Bax, Bcl-2 (B); caspase-3, cleaved caspase-3 (G); and NF- κ B (H). The column chart shows antibody relative intensity of COX-2 (C); iNOS (D); Bax (E); Bcl-2(F); cleaved caspase-3 (I); NF- κ B (J). The protein expression was examined by the western blotting analysis in brain tissues from all experimental groups. * $p < 0.05$, ** $p < 0.01$ vs. control group; # $p < 0.05$, ## $p < 0.01$ vs. model group.



fibrauretime-treated group occurred for the donepezil-treated group.

3.5 Effects of fibrauretime on inflammation of the brain in D-galactose/ AlCl_3 -treated mice

The previous study has confirmed that AD is related to the release of pro-inflammatory cytokines, including tumour necrosis factor- α (TNF- α), interleukin-1 β (IL-1 β) and interleukin-33 (IL-33).^{37–40} The brain tissue levels of TNF- α and IL-1 β were determined by the ELISA assay. As indicated in Fig. 5A and B, a single D-galactose/ AlCl_3 administration resulted in obviously higher levels of TNF- α and IL-1 β in the brain, compared with the control group ($p < 0.01$). However, these increases were reversed after fibrauretime administration in a dose-dependent manner ($p < 0.05$ or $p < 0.01$). The levels of TNF- α and IL-1 β in the donepezil-administered mice were also significantly reduced ($p < 0.01$). The levels of IL-33 and its receptor ST2 were detected by IHC analysis. As shown in Fig. 5C and D, the positive expression area of IL-33 and ST2 levels of the model group was clearly increased, compared with the control group. But fibrauretime treatment resulted in dose-dependent reduction. And the administration of donepezil also reduced the levels of IL-33 and ST2 ($p < 0.05$). Meanwhile, to further discuss the mechanisms underlying the advantageous effects of fibrauretime on D-galactose/ AlCl_3 -induced neuroinflammation, the levels of NF- κ B, COX-2 and iNOS were detected in all groups (Fig. 5E–G and 6A and H). Compared with the control group, the positive expression area of these protein levels of the model group was clearly improved. However, these increases were attenuated by fibrauretime at doses of 80, 160 and 320 mg kg^{-1} ($p < 0.05$). So did the donepezil-treated group ($p < 0.05$).

3.6 Effects of fibrauretime on apoptosis of the brain in D-galactose/ AlCl_3 -treated mice

To measure the degree of apoptosis in the brains, the protein expressions of the cleaved caspase-3, anti-apoptotic factor Bcl-2 and pro-apoptotic factor Bax in all experimental groups of mice were detected. In the immunohistochemical analysis, as shown in Fig. 5H and I, the hippocampal pyramidal cells are stained in brown, and the positive cells are stained in yellow. Compared with the control group, the rates of positive expressions of Bax located in hippocampal pyramidal cytoplasm were observed to be clearly higher in the model group ($p < 0.01$). There is a clear reduction effect in the fibrauretime administration group ($p < 0.05$). The donepezil-treated group showed the same therapeutic effect ($p < 0.05$). Conversely, the positive expression of Bcl-2 in the hippocampal pyramidal cytoplasm has increased in the fibrauretime groups compared to the model group ($p < 0.05$). So did the donepezil-treated group ($p < 0.05$). The results of western blotting showed that D-galactose/ AlCl_3 exposure significantly increased the expression levels of cleaved caspase-3 and Bax and reduced the expression level of Bcl-2 compared with the control group ($p < 0.01$). However, similar to the results of immunohistochemistry, the levels of Bax, cleaved caspase-3 of the fibrauretime-treated groups significantly reduced ($p <$

0.01). So did the donepezil-treated group ($p < 0.05$). (Fig. 6D–F) ($p < 0.05$).

The TUNEL assay can be used to detect DNA strand breaks that are associated with apoptosis.⁴¹ To further confirm whether fibrauretime treatment could inhibit nerve cell apoptosis in D-galactose/ AlCl_3 -induced AD, the TUNEL assay was conducted (Fig. 4B). The effect on hippocampal neuron cells was represented by the colour stained, and positive staining was in brown. The results showed that fibrauretime pretreatment with 80 mg kg^{-1} , 160 mg kg^{-1} and 320 mg kg^{-1} decreased the number of D-galactose/ AlCl_3 -induced TUNEL-positive cells ($p < 0.05$ or $p < 0.01$); the same outcome occurred for the donepezil group. These results showed that fibrauretime showed suppressive effects against D-galactose/ AlCl_3 -induced hippocampal neuron cell apoptosis.

4 Discussion

In the present study, a D-galactose/ AlCl_3 -induced AD mice model was established to investigate the anti-AD activities of fibrauretime by intragastrical administration. Our results showed that fibrauretime significantly enhanced the spatial short-term working memory in AD mice in the Y-maze test and the spatial learning and memory in the Morris water maze. The levels of A β , P-Tau, Tau and AchE in brain tissue were evidently increased in the model group and were reversed by fibrauretime. In contrast, a significant increase in the levels of Ach and ChAT was observed in the fibrauretime-treated groups compared with the model group. The results indicated that fibrauretime exhibited ameliorative effects on D-galactose/ AlCl_3 -induced AD in mice, in a dose-dependent manner.

A great deal of evidence has implicated oxidative stress and free radical damage in the pathogenesis of AD. The damage found in AD includes advanced glycation end products,⁴² nitration,⁴³ lipid peroxidation adduction products,^{44,45} and carbonyl-modified neurofilament protein and free carbonyls.^{46,47} The antioxidant factors are represented by anti-oxidant enzymes such as superoxide dismutase (SOD), glutathione peroxidase (GPX), catalase or aldehyde dehydrogenase, and non-enzymatic antioxidant factors. The reaction of the reduction of free radicals is catalysed by the antioxidant enzymes, diminishing their power and hence oxidative cytotoxicity.⁴⁸ ELISA results showed that neuronal oxidative stress was induced by D-galactose/ AlCl_3 in the model group, evidenced by increased malondialdehyde (MDA) and nitric oxide (NO) levels and a decline in glutathione (GSH), catalase (CAT) and superoxide dismutase (SOD) activity. All of the above abnormality was reduced by fibrauretime in the present work. As described above, our findings illustrated which fibrauretime pretreatment recovered antioxidant ability by suppressing oxidative stress.

Numerous studies demonstrated that neuroinflammation plays a critical role in the pathogenesis of AD, showing a clear involvement of several inflammatory pathways.⁴⁹ NF- κ B is a pro-inflammatory transcription factor located in the cytoplasm *via* binding to the inhibitory I κ B protein. I κ B is degraded by the proteasome when it is stimulated by pathogens. Then, NF- κ B is



freed to translocate from the cytoplasm to the nucleus where it promotes the transcription of target genes such as TNF- α and IL-1 β . At the same time, the up-regulation of iNOS and COX-2, *via* the NF- κ B pathway, is another central mechanism in inflammatory response processes.⁵⁰ Growing evidence shows that iNOS and COX-2 are clearly highly expressed at sites of inflammation. Moreover, iNOS and COX-2 have synergistic effects to accelerate the inflammatory response.⁵¹ IL-33 serves as a proinflammatory cytokine and an intra-cellular nuclear factor regulating transcription. IL-33 may be a specific pro-inflammatory cytokine in the CNS.⁵² IL-33 receptors containing a trans-membrane portion called ST2 are expressed in microglia and astrocytes. The glial cells respond by proliferating and releasing TNF- α , IL-1 β , and NO.⁴⁰ IL-1 β and TNF- α are released by IL-33 from inflammatory cells.³⁹ Both IL-1 β and TNF- α were increased in the brains of AD patients compared to control subjects.³⁸ Fibrauretime has traditionally been used as the anti-inflammatory agent. However, its anti-inflammatory mechanism is not clear. In this study, to discuss the potential molecular mechanisms and the anti-AD effects of fibrauretime against D-galactose/AlCl₃-induced inflammation in depth, we detected the expression levels of NF- κ B, COX-2, iNOS, IL-33 and ST2. Western blotting and immunohistochemistry analysis of the results showed that fibrauretime pretreatment significantly down-regulated the increase in expression levels of NF- κ B, COX-2, iNOS, as well as IL-33 and ST2 after D-galactose/AlCl₃ exposure. Moreover, ELISA results revealed that D-galactose/AlCl₃ administration increased the levels in TNF- α and IL-1 β . Nevertheless, the increase in inflammatory cytokine levels against D-galactose/AlCl₃-induced mice was dramatically suppressed by fibrauretime. The possible molecular mechanisms are involved in the suppression of such cytokines by fibrauretime. In short, the results suggested that fibrauretime could act as an anti-inflammatory agent against D-galactose/AlCl₃-induced AD.

There is growing evidence that AD is related to apoptosis. The accumulation of A β can induce neuronal apoptosis, which plays a vital role in AD pathogenesis.⁵³ The conditions inducing apoptosis that include reactive oxygen species (ROS), nitric oxide (NO), glucocorticoids and Bax overexpression are known as major factors causing the release of cytochrome *c* (Cyt *c*).^{54,55} The Bcl-2 family of proteins strictly regulates Cyt *c* release.⁵⁶ The family is composed of anti-apoptosis protein such as Bcl-2 and pro-apoptotic protein such as Bax. The balance between anti- and pro-apoptotic proteins determines the activation of intrinsic apoptotic signals involving mitochondria.⁵⁷ When upstream caspases such as caspase-9 are activated, downstream effector caspase cleavage such as caspase-3 will be triggered, and they can cleave cytoskeletal and nuclear proteins to induce apoptosis.⁵⁸ For evaluated levels of apoptosis in brain tissues following D-galactose/AlCl₃ administration, the effects of fibrauretime were assessed by cleaved caspase-3, pro-apoptotic factor Bax and anti-apoptotic factor Bcl-2 in every pretreatment group. The findings from protein detection analyses of brains markedly showed that the expression levels of Bax, caspase-3, and cleaved caspase-3 were suppressed, while the expression level of Bcl-2 was relatively activated. Additionally, the results of TUNEL staining displayed that the apoptosis rate

in brain tissues was clearly reduced by fibrauretime when compared with the model group. In summary, the above results demonstrated that fibrauretime inhibited the apoptosis of brain tissues in D-galactose/AlCl₃-induced AD mice.

In summary, AD is a complex disease involving many genetic and environmental factors. Oxidative stress has been shown to induce apoptosis by inducing cytochrome *c* release and activating caspase-9 and caspase-3,⁵⁹ and overproduction of free radicals would be followed by an inflammatory response.⁶⁰ Among various potential causes, inflammation has been suggested to be a predisposing factor in the development of neurodegenerative diseases including AD.

Nuclear factor kappa B (NF- κ B) is a key inflammatory mediator that regulates the expression of various genes that involved in apoptosis and inflammation.⁶¹ NF- κ B is usually kept inactive in the cytoplasm through association with an endogenous inhibitor protein of the I κ B family.⁶² Activated NF- κ B could increase the expression of other inflammatory mediators such as COX-2, iNOS, IL-1 β and TNF- α , which might be triggered memory impairment.²¹ Interestingly, ROS accumulation in response to TNF- α is regulated in an NF- κ B-dependent manner.⁶³ Previous study has found that inhibition of NF- κ B inhibits apoptosis during the resolution of inflammation *in vivo* and expression of genes involved in apoptosis, including the pro-apoptotic Bcl-2 homolog Bax, was modulated by NF- κ B inhibitors.⁶⁴ Thus it can be seen that NF- κ B plays a very important role and may be a key factor in the onset of AD. The immunohistochemistry and western blotting results from our study demonstrated that fibrauretime can inhibit the activation of NF- κ B. Several studies have also revealed that fibrauretime can restrain the activation of NF- κ B in other cell models or animal models.⁶⁵⁻⁶⁹

So fibrauretime from *Fibraurea recisa* Pierre ameliorate D-galactose/AlCl₃-mediated Alzheimer's disease probably by suppressing the NF- κ B pathways.

5 Conclusion

In addition to being clinically used as an anti-inflammatory drug, fibrauretime has been found to have a significant anti-AD effect, the mechanism is inextricably linked to anti-inflammatory, anti-oxidation and anti-apoptosis.

Conflicts of interest

There are no conflicts of interest to declare.

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