Original Article

Polygalasaponin XXXII, a triterpenoid saponin from *Polygalae Radix*, attenuates scopolamine-induced cognitive impairments in mice

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Aim: Recent studies show that the extract of a Chinese herb *Polygalae Radix* exerts cognition-enhancing actions in rats and humans. The aim of this study was to characterize the pharmacological profiles of active compounds extracted from *Polygalae Radix*.

Methods: Two fractions P3 and P6 and two compounds PTM-15 and polygalasaponin XXXII (PGS32) were prepared. Neuroprotective effects were evaluated in primary cortical neurons exposed to high concentration glutamate, serum deficiency or H₂O₂. Anti-dementia actions were assessed in scopolamine-induced amnesia in mice using step-through avoidance tests and channel water maze tests. After conducting the channel water maze tests, TrkB phosphorylation in mouse hippocampus was detected using Western blotting. Long-term potentiation (LTP) was induced in the dentate gyrus in adult rats; PGS32 (5 μ L 400 μ mol/L) was injected into the lateral cerebral ventricle 20 min after high frequency stimulation (HFS).

Results: Compared to the fraction P6, the fraction P3 showed more prominent neuroprotective effects *in vitro* and cognition-enhancing effects in scopolamine-induced amnesia in mice. One active compound PGS32 in the fraction P3 exerted potent cognition-enhancing action: oral administration of PGS32 (0.125 mg kg⁻¹ d⁻¹) for 19 days abolished scopolamine-induced memory impairment in mice. Furthermore, PGS32 (0.5 and 2 mg kg⁻¹ d⁻¹) significantly stimulated the phosphorylation of TrkB in the hippocampus. Intracerebroventricular injection of PGS32 significantly enhanced HFS-induced LTP in the dentate gyrus of rats.

Conclusion: PGS32 attenuates scopolamine-induced cognitive impairments in mice, suggesting that it has a potential for the treatment of cognitive dysfunction and dementia.

Keywords: *Polygalae Radix*; polygalasaponin XXXII; triterpenoid saponin; neuroprotection; scopolamine; dementia; tyrosine kinase B; LTP

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Introduction

Neurodegenerative diseases are a group of chronic, progressive disorders characterized by the gradual loss of neurons in discrete areas of the central nervous system^[1]. These groups of diseases include highly prevalent diseases, such as Parkinson's disease and Alzheimer's disease, and other rare diseases, including prion diseases (also known as transmissible spongiform encephalopathies), spinocerebellar ataxia, amyotrophic

lateral sclerosis and Huntington's disease^[2,3]. As age-related diseases, neurodegenerative diseases are becoming an increasing burden as the human population ages, with approximately 5.1 million households affected by Alzheimer's disease, and almost 300 000 people of age 65 years or older afflicted with Parkinson's disease in the US as of 2010^[4].

Traditional therapies for neurodegenerative diseases enhance cognition and improve the quality of life of patients, but they do not alter the natural course or the ultimate outcome of these diseases^[5]. Using herbal treatments in neurodegenerative disease therapies has generated unprecedented levels of interest because of the multi-function, multi-target

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characteristics of these treatments^[6]. Many researchers have developed new compounds or active extracts from herbal materials for the treatment of neurodegenerative disorders.

Polygalae Radix, a well-known herb listed in the Chinese Pharmacopoeia (2015 edition), has been prescribed as a tonic, antipsychotic, expectorant and detumescent for thousands of years in traditional Chinese medicine. Modern pharmacological studies have already demonstrated that Polygalae Radix extract exhibits a wide variety of effects, such as hypotensive^[7], anti-inflammatory^[8], anti-depressive^[9], anti-oxidative and anti-apoptotic^[10] effects, cognitive improvements and cerebral protection^[11]. The Polygalae Radix extract BT-11 has been shown to have memory-enhancing effects in rats with scopolamine-induced amnesia^[12] and cognition-improving effects in two randomized, double-blind, placebo-controlled comparison studies, one in healthy adults and one in elderly humans^[13, 14]. However, the biologically active components of Polygalae Radix extract remains unclear, and additional preclinical studies are necessary to develop more effective monomers from Polygalae Radix extracts.

In the present study, we attempted to screen for polygala saponin monomers with neuroprotective and anti-dementia effects and to further clarify our understanding of their underlying mechanisms.

Materials and methods

Drugs and reagents

Glutamate, nerve growth factor (NGF, purity >97%), MTT, huperzine A (Hup-A), poly-D-lysine (PLL), and cytarabine (Ara-C) were purchased from Sigma Chemical Company (Sigma-Aldrich (St Louis, MO, USA)). Scopolamine hydrobromide was purchased from the National Institute for the Control of Pharmaceutical and Biological Products (Beijing, China). Donepezil was provided by the Department of Chemosynthesis, Institute of Materia Medica, Chinese Academy of Medical Sciences and Peking Union Medical College (Beijing, China). Dulbecco's modified Eagle's medium/nutrient mixture F-12 (DMEM/F12), Dulbecco's modified Eagle's medium (DMEM), fetal bovine serum (FBS), and horse serum (ES) were purchased from Gibco BRL (New York, NY, USA). Anti tyrosine kinase B (TrkB) and phosphorylated TrkB (p-TrkB) antibodies were obtained from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Other materials were purchased from other commercial sources.

Compound preparation

The compounds were prepared as previously described^[15]. Fraction Fc-2 and fraction Fc-3 were P3, while Fraction Fc-7 was P6. Fraction Fc-2 was separated by preparative medium pressure liquid chromatography and eluted with MeOH-H₂O to obtain nine fractions (Fc2-1 to Fc2-9), and then PTM-15 and polygala saponin XXXII (PGS32) were further separated from Fc2-8 by semi-preparative high pressure liquid chromatography (MeOH-H₂O 70:30+0.05% CF₃COOH, 7 mL/min). PGS32 was first isolated from the root of *Polygala japonica* Houtt by Zhang *et al* in 1996^[16].

Primary cortical neuron culture and treatment

Primary cortical neurons were prepared from rat embryos at 17±19 d gestation as previously described^[17]. Cerebral neocortices were removed and cultured in DMEM/F12 containing 10% ES and 10% FBS. Cells were plated in PLL-coated 96-well plates at 1×10⁶/mL for toxicity experiments. Then, plates were incubated at 37°C in a humidified atmosphere with 5% CO₂. Two days later, 10 µg/mL Ara-C was added to the cultures to inhibit the outgrowth of various cells for 48 h, including glial cells, mechanocytes, and nerve stem cells. The primary cortical neurons were cultured for another seven days, and they were then used for treatment. The media were changed every two days. Nerve growth factor was used as the positive control.

Glutamate at a concentration of 10 mmol/L was used as an inducer for neurotoxicity in primary cortical neurons. The test compounds were added to each of the treated groups at concentrations of 1, 10, and 100 µg/mL. Twenty-four hours after treatment, 10 µL MTT (5 mg/mL) was added to each well, and cells were cultured in the incubator for 4 h. Then, 100 µL solubilization solution [10% sodium dodecyl sulfate (SDS) in 0.01 mol/L HCl] was added to each well of the 96-well plate. The plate was incubated overnight at 37°C, and the optical absorbance of the cells was measured at 570 nm with a microtiter plate reader.

We conducted MTT assays to measure the survival rate of primary cortical neurons injured by serum deficiency. The original culture media were replaced by media containing 0.5% FBS and 0.5% ES, with or without Polygalae Radix-derived compounds. The cells in the control group were treated with complete culture media but no compounds. The cells were cultured for another 24 h. Then, an MTT assay was performed to measure the survival rate of primary cortical neurons.

PC12 cell culture and treatment

PC12 cells were cultured in DMEM supplemented with 5% ES and 5% FBS. The media were changed every 2 days, and cells were seeded on PLL-coated plates at 5×10⁴/mL. After 24 h subculture, cells were transferred to culture media containing 100 µmol/L H₂O₂ with or without Polygalae Radix-derived compounds. After 24 h, the MTT assay was performed to measure the survival rates of PC12 cells.

Animals and drug treatment

Male Kunming mice (18–20 g), C57BL/6J mice (18–20 g) and Wistar rats (230–260 g) were provided by the Experimental Animal Center of the Chinese Academy of Medical Sciences. They were housed in a room under temperature and light control (23°C, 12 h-light cycle) and had free access to food and water. All animal experiments were performed in accordance with the National Institutes of Health Guide for the Care and Use of Laboratory Animals (NIH Publications No 80–23), and the animal study was approved by the Animal Care Committee of the Chinese Academy of Medical Sciences and the Peking Union Medical College.

Eighty-four Kunming mice were divided into 7 groups as follows: control group, model group, 7.5 mg/kg P3 group, 30 mg/kg P3 group, 7.5 mg/kg P6 group, 30 mg/kg P6 group, and 5 mg/kg donepezil group (the last of which was the positive control group). The experiment lasted for 16 days. P3, P6 and donepezil were suspended in double-distilled water (DDW) and orally administered (ig) to the mice (0.1 mL/10 g) every day. The same volume of DDW was given to the control group and the model group. During the behavioral tests from the 10th day onward, the P3, P6, donepezil and model groups were injected with scopolamine (5 mg/kg d 10–d 11 for step-through avoidance tests, 2 mg/kg d 13–d 16 for channel water maze tests, ip, suspended in saline) 30 min prior to the test, whereas the control group was injected with the same volume of normal saline.

Eighty-four C57BL/6J mice were divided into seven groups as follows: control group, model group, PTM-15 5 mg/kg group, PTM-15 20 mg/kg group, PGS32 5 mg/kg group, PGS32 20 mg/kg group and positive control group. PTM-15 and PGS32 were suspended in DDW and administered orally (ig) to mice at doses of 5 or 20 mg/kg. Donepezil (5 mg/kg) was suspended in DDW and orally administered (ig) to the positive control group. The same volumes (0.1 mL/10 g) of DDW were given to the control group and model group. The experiment lasted for 13 days. The PTM-15, PGS32, donepezil and model groups were injected with scopolamine (2 mg/kg d 10–d 13, ip, suspended in saline) 30 min prior to the channel water maze test, whereas the control group was injected with the same volume of normal saline.

Seventy-two C57BL/6J mice were divided into six groups as follows: control group, model group, PGS32 groups (0.125, 0.5, or 2 mg/kg), and positive control group. PGS32 at different concentrations were suspended in DDW and administered orally (ig) to mice. Hup-A (0.05 mg/kg) was suspended in DDW and orally administered (ig) to the positive control group. The same volume (0.1 mL/10 g) of DDW was given to the control and model groups. The experiment lasted for 19 days. The PGS32, Hup-A, and model groups were injected with scopolamine (2 mg/kg, d 16–d 19, ip, suspended in saline) 30 min prior to the channel water maze test, whereas the control group was injected with the same volume of normal saline.

Step-through avoidance test

The step-through avoidance tests were conducted following procedures described previously with minor modification^[18]. The apparatus contained a chamber with two compartments (dark/light), which were connected by a guillotine door. The dark compartment contained a stainless steel grid floor that could deliver electric shocks. During the acquisition trial, each mouse was placed in the light compartment. After 180 s of habituation, the guillotine door was raised, and the initial latency time of each mouse to enter the dark compartment was recorded. Immediately after the mouse entered the dark compartment, the guillotine door was closed, and an electric shock was delivered. Five seconds later, the mouse was removed

and returned to its home cage. Twenty-four hours after training, the latency time was measured following the same procedures in the acquisition trial, but no electric shock was delivered. The number of times that the mouse entered the dark side was recorded as the number of errors. The latency time to enter the dark compartment was recorded up to 300 s.

Channel water maze test

The channel water maze test was performed as previously described with minor modifications^[19]. A channel water maze (73 cm×42 cm×30 cm) made of black glass was designed by the Pharmacological Institute of the Chinese Academy of Medical Sciences. The tank was filled with water to a depth of 20 cm and maintained at 22±2°C. There were four impasses and a platform at the terminal in the water maze. Mice could avoid drowning by climbing onto the platform.

On the first day of training, starting point A was cut off by a black bar. Each mouse was placed on the platform at the terminal for 15 s, and then it was placed on A with its head facing the bar. The mouse was allowed to swim freely twice for 2 min during each training session. The latency time to find the platform and the number of errors in which the mouse swam into impasse 1 were recorded. Any mouse that failed to find the platform after 2 min was guided to the platform, where it was placed for 15 s. On the second day of training, starting point B was cut off, and each mouse was placed on B; the latency time to find the platform and the number of errors in which the mouse swam into impasse 1, 2, or 3 were recorded. On the third day of training, each mouse was placed on C. The latency time to find the platform and the number of errors in which the mouse swam into impasse 1, 2, 3, or 4 were recorded. The fourth day was the testing day. Each mouse was placed on C; the latency time to find the platform and the number of errors in which the mouse swam into impasse 1, 2, 3, or 4 were recorded.

Western blotting

After channel water maze testing, six mice from each group were decapitated, and hippocampus samples from the mice were homogenized thoroughly and then lysed in RIPA lysis buffer [50 mmol/L Tris (pH 7.4), 150 mmol/L NaCl, 1% NP40, 0.5% sodium deoxycholate and 0.1% SDS]. Protein concentrations were measured with a BCA kit (Applygen Technologies, Beijing, China). The sample lysates were solubilized in SDS sample buffer, separated by SDS-PAGE, and transferred to a PVDF membrane as previously described^[20]. The membrane was blocked with 3% BSA and incubated with primary antibody, followed by horseradish peroxidase-conjugated secondary antibody. Detection was performed using an enhanced ECL plus detection system. The relative density of the bands was analyzed by Gelpro 32 software.

Electrophysiological assays

Twelve Wistar rats (230–260 g), 3 months old, were randomly assigned to either the control group ($n=6$) or the PGS32 group ($n=6$). Recordings of evoked potentials, intracerebroven-

tricular (icv) drug delivery, data collection and analysis were made as previously described^[21]. In brief, the amplitude of the population spike (PS) was employed to assess the excitation level of the granular cell population in the dentate gyrus. The baseline was recorded for 30 min before long-term potentiation (LTP) was induced by high frequency stimulation [HFS: 10 bursts of 5 stimuli (at 100 Hz, 0.2 ms duration, 200 ms inter-burst interval)]. Twenty minutes after HFS, 5 μ L 400 μ mol/L PGS32 (icv, final concentration of 1 μ mol/L) was injected into each test subject, while the same volume 0.4% DMSO-Normal Saline was injected into each control rat, and the intensity of LTP was recorded for another hour.

Statistical analyses

Data are expressed as the mean $\pm$ SEM as indicated. The data were analyzed using Graph Pad Prism 5.0 (San Diego, CA, USA) and the SPSS 16.0 software (Chicago, IL, USA). Group differences in PS amplitude in the electrophysiological assays were analyzed using two-way analysis of variance (ANOVA) with repeated measures. Group differences in cytotoxicity assays and cognitive behavior tests were evaluated using one-way ANOVA followed by Duncan's multiple-range test. The accepted level of significance was set at $P<0.05$.

Results

Neuroprotective effects of P3 and P6 in *in vitro* models

Figure 1A illustrates that treatment with 10 mmol/L glutamate resulted in a decrease in cell viability. P3 and P6 protected against the effects of glutamate on primary cortical neurons. P3 (1, 10, 100 μ g/mL) significantly increased the cell survival rate ($P<0.01$, $P<0.05$, $P<0.01$), while P6 showed neuroprotective effects only at the concentration of 100 μ g/mL ($P<0.01$). As shown in Figure 1B, P3 significantly increased the survival rate of primary cortical neurons under serum deficient conditions and did so in a dose-dependent manner (1 μ g/mL, $P<0.05$; 10 μ g/mL, $P<0.01$; 100 μ g/mL, $P<0.01$). Figure 1C showed that the viability of PC12 cells exposed to 100 μ mol/L H_2O_2 for 24 h was less than the viability in the normal group ($P<0.01$), and P3 showed neuroprotective effects against H_2O_2 -induced PC12 cell death at a concentration of 10 μ g/mL ($P<0.01$).

Effects of P3 and P6 on learning and memory performance in a scopolamine-induced amnesia model in mice

Increased latency times may serve as indicators of memory enhancement, whereas decreased latency times indicate memory impairment in the step-through avoidance test. No significant differences in step-through latency time were observed between groups during the acquisition trials (data not shown). During the retention trials, the control group mice hesitated to re-enter the dark compartment, but the model group mice re-entered the dark compartment more frequently ($P<0.01$). The model group mice showed shorter latency times ($P<0.05$). In both the 7.5 and the 30 mg/kg P3-treated groups, mice showed longer latency times and fewer error trials, while P6 only reversed scopolamine-induced memory impairments

Figure 1. (A) Effects of P3 and P6 on the survival rate of primary cortical neurons injured by glutamate. ** $P<0.01$ vs Ctrl. # $P<0.05$, ## $P<0.01$ vs Glu. (B) Effects of P3 on the survival rate of primary cortical neurons injured by serum deficiency. ** $P<0.01$ vs Ctrl. # $P<0.05$, ## $P<0.01$ vs serum deficiency. (C) Effects of P3 on the survival rate of PC12 cells injured by H_2O_2 . Mean $\pm$ SEM. $n=9-11$. ** $P<0.01$ vs Ctrl. ## $P<0.01$ vs 100 μ mol/L H_2O_2 .

at a dose of 7.5 mg/kg. Additionally, similar patterns were observed following administration of 5 mg/kg donepezil, an acetylcholinesterase inhibitor that was used as a positive control (Figure 2A and 2B).

In the channel water maze test, the latency time for P3 and P6 treated mice to reach the platform significantly decreased compared with the mice in the model group, and the mice treated with P3 7.5 mg/kg had the best performance (Figure 2C). The number of error trials in the model group was higher than in the control group ($P<0.01$). The total numbers of errors declined significantly only in the P3-treated group (Figure 2D). These results suggest that P3 showed better neuroprotective and cognition-enhancing effects than did P6. Hence, we chose

Figure 2. (A) Effects of P3 and P6 on the latency time tested 24 h after training in the step-through passive avoidance test. Mean±SEM. $n=11-12$. (B) Effects of P3 and P6 on the error numbers tested 24 h after training in the step-through passive avoidance test. Mean±SEM. $n=11-12$. (C) Effects of P3 and P6 on the latency time after training in the channel water maze test. Mean±SEM. $n=9-11$. (D) Effects of P3 and P6 on the error numbers tested 24 h after training in the channel water maze test. Mean±SEM. $n=9-11$. * $P<0.05$, ** $P<0.01$ vs Ctrl. # $P<0.05$, ## $P<0.01$ vs Model.

P3 for our further research.

Effects of PTM-15 and PGS32 on learning and memory performance in the scopolamine-induced amnesia model in mice

PTM-15 and PGS32 were further separated from P3. In the channel water maze test, we observed that the latency time and number of errors in mice treated by the administration of scopolamine (1 mg/kg) increased significantly compared to those of vehicle-treated control mice ($P<0.01$, $P<0.01$). Furthermore, the increased latency times and error numbers induced by the administration of scopolamine were significantly reversed by treatment with 5 mg/kg PGS32 (Figure 3). These results demonstrated that PGS32 attenuated scopolamine-induced learning and memory impairments in mice. In our behavioral test, the 20 mg/kg PGS32 treatment was not as effective as the 5 mg/kg treatment; these results suggested that a lower dose of PGS32 might be more effective. Thus, we designed a new experiment with a concentration gradient of 0.125, 0.5, and 2 mg/kg.

Effects of PGS32 on learning and memory performance in the scopolamine-induced amnesia model in mice

In the channel water maze test, we observed that the latency times and error numbers in mice treated with scopolamine (2 mg/kg) increased significantly compared to the vehicle-treated control mice ($P<0.05$, $P<0.05$). The latency times to reach the platform of PGS32 treated mice decreased signifi-

cantly compared with the model group, and the total error numbers declined significantly in the PGS32-treated group. PGS32 (0.125, 0.5, and 2 mg/kg) treatment reversed the memory impairments induced by scopolamine. Furthermore, Hup-A treatment showed similar cognition-improving effects (Figure 4).

Effects of PGS32 on TrkB and p-TrkB in the hippocampal lysates of mice following the channel water maze test

The activation of TrkB by BDNF mediates intracellular signal transduction in which the phosphorylation of TrkB plays a pivotal role. Western blotting results showed that the level of p-TrkB was significantly decreased in the scopolamine-treated mice compared with the control group, whereas there was no significant difference in total TrkB expression between the groups. PGS32 (0.5 or 2 mg/kg) and Hup-A treatment significantly up-regulated p-TrkB compared with the model group (Figure 5).

Effects of PGS32 on long-term potentiation induced by high-frequency stimulation in the dentate gyrus of anesthetized rats

LTP was induced by high frequency stimulation after a stable baseline of 20 min. Before treatment, there were no significant differences in HFS-induced PS amplitudes between these two groups (Control: $273\pm40\%$; PGS32: $283\pm32\%$). PGS32 or vehicle (DMSO) was injected into the lateral cerebral ventricle of rats 20 min after HFS. Compared with the control group,

Figure 3. Effects of PTM-15 and PGS32 on scopolamine-induced learning and memory deficits in the channel water maze test. (A) Effects of PTM-15 and PGS32 on the latency time tested 24 h after training in the channel water maze test. (B) Effects of PTM-15 and PGS32 on the error numbers tested 24 h after training in the channel water maze test. Mean $\pm$ SEM. $n=10-12$. ** $P<0.01$ vs Ctrl. ## $P<0.01$ vs Model.

the PS amplitude of the PGS32 treatment group was enhanced significantly ($P<0.01$). PS amplitudes increased to $310\%\pm 20\%$ within 20 min after PGS32 application, and at 40 min, 50 min, 55 min after PGS32 treatment, PS amplitudes increased to $340\%\pm 24\%$, $378\%\pm 27\%$, and $366\%\pm 26\%$, respectively. These results suggest that PGS32 significantly enhanced hippocampal LTP. Therefore, PGS32 may promote synaptic plasticity in the hippocampus (Figure 6).

Discussion

In the current study, we demonstrated that P3 not only has neuroprotective effects on excitotoxicity, oxidative stress and serum deficiency induced damage but also has cognition-improving effects in scopolamine-induced amnesia model mice. Moreover, we separated P3 to yield PGS32, and we discovered that PGS32 reversed scopolamine-induced amnesia in mice at a dose of 0.125 mg/kg. These results suggest that PGS32 might be one of the active monomers in *Polygalae Radix*.

Glutamate and ROS can contribute to neuronal damage^[22]. Compounds ameliorating these types of damage may be effective in treating neurodegenerative diseases^[23]. We used MTT assays to screen for neuroprotective compounds. We discovered that P3 protected primary cortical neurons against serum deficiency and glutamate-induced damage and PC12 cells

Figure 4. Effects of PGS32 (0.125–2 mg/kg) on scopolamine-induced learning and memory deficits in the channel water maze test. (A) Effects of PGS32 on the latency time tested 24 h after training in the channel water maze test. (B) Effects of PGS32 on the error times tested 24 h after training in the channel water maze test. Mean $\pm$ SEM. $n=10-12$. * $P<0.05$ vs Ctrl. # $P<0.05$, ## $P<0.01$ vs Model.

against H_2O_2 -induced death. Several researchers have demonstrated that *Polygalae Radix* extract provides neuroprotection of PC12 cells against 6-hydroxydopamine (6-OHDA)-induced damage and of primary cortical neurons against $A\beta_{25-35}$ -induced cell damage^[24, 25]. These results suggest that P3 may be effective in the treatment of various neurodegenerative diseases.

Learning and memory impairments are primary clinical symptoms observed in all types of dementia^[26]. We used a scopolamine-induced amnesia model to further investigate the efficacy of P3. As a muscarinic receptor antagonist, scopolamine causes profound memory impairments by antagonizing muscarinic cholinergic receptors in the central cholinergic systems^[27]. Donepezil is a selective reversible inhibitor of acetylcholinesterase; Hup-A, an alkaloid isolated from the Chinese folk medicine *Huperzia serrata*, is a reversible and selective inhibitor of AChE^[5, 28]. These improve cognitive deficits in a broad range of animal models and have been used in treating Alzheimer's disease in the clinic. Avoidance step-through tests and channel water maze tests were used to eval-

Figure 5. Effects of PGS32 (0.125–2 mg/kg) on the levels of phosphorylated TrkB and TrkB in the hippocampal lysates of mice following the channel water maze test. (A) Representative immunoblots of the hippocampal lysates of the mice. (B) The ratio of p-TrkB/t-TrkB after PGS32 treatment. Mean $\pm$ SEM. $n=6$. ** $P<0.01$ vs Ctrl. ## $P<0.01$ vs model.

Figure 6. Effects of PGS32 on long-term potentiation induced by 10 bursts of 5 stimuli (100 Hz, 0.2 ms stimulus duration, 200 ms interburst interval) in the dentate gyrus of anesthetized rats. The baseline was recorded for 30 min before HFS. Twenty minutes after HFS-induced LTP, PGS32 1 μ mol/L or 0.4% icy dimethyl sulfoxide (DMSO) was administered intracerebroventricularly, and the intensity of LTP was recorded for another hour. Mean $\pm$ SEM. $n=6$. ** $P<0.01$ vs Ctrl.

uate learning and memory ability in rodents. We discovered that P3 showed cognition-enhancing effects in scopolamine-induced amnesia model in mice. To accelerate these developments, we separated polygala saponin monomers from P3. Of these polygala saponin monomers, PGS32 (1 μ mol/L) showed neuroprotective effects against serum deficiency and glutamate induced damage in PC12 cells^[15]; PGS32 (0.125, 0.5, and 2 mg/kg) treatments reversed the learning and memory impairments induced by scopolamine. We focused on PGS32

because it was more potent *in vivo* and *in vitro*, and its underlying mechanisms are discussed below.

Polygalae radix extract can protect against *N*-methyl *D*-aspartate (NMDA) neurotoxicity^[29] and induce the expression of brain-derived neurotrophic factor (BDNF)^[30], which is involved in the modulation of synaptic transmission and plasticity^[31]. Activation of TrkB by BDNF results in the phosphorylation of several tyrosine residues and subsequently the stimulation of intracellular signal transduction pathways, such as the MAPK, PI3K and phospholipase C- γ pathways^[32]. Phosphorylation of TrkB is associated with the recruitment of pro-survival signaling pathways that lead to long-term neuroprotection and improved cognition^[33]. We have demonstrated previously that PGS32 can activate ERK and CREB and further increase the expression of BDNF^[34]. In this study, we found that PGS32 up-regulated the phosphorylation of TrkB. It was reported that compared with the vehicle control rats, p-TrkB was significantly decreased in the scopolamine-infused rats^[35]. Galantamine, a cholinergic stimulator used as a positive control, was shown to restore p-TrkB levels and improved cognitive functioning in AD-like rodent models in another study, which is inconsistent with our results. These results suggest that PGS32 may improve learning and memory by enhancing the BDNF/TrkB/MAPK cascade.

BDNF/TrkB signaling plays a fundamental role in both developmental and adult synaptic plasticity^[36]. LTP of synaptic transmission in the hippocampus is the primary experimental model for investigating the synaptic basis of learning and memory in vertebrates^[37]. Electrophysiological LTP assays are widely used in screening candidate cognition-improving drugs. Learning and memory can be studied at many levels, from the molecular to the behavioral. For our cognitive function test, which was a relatively long-term experiment, we chose oral administration to mimic the practical application of pharmaceutical preparations in the clinic. Oral administration is a mild and easy route for drug delivery. For the LTP measurement, we focused on the immediate impact of PGS32 on basic synaptic transmission to investigate the underlying mechanisms of its cognition improving effects, and we chose intracerebroventricular administration. Our previous studies have shown that PGS32 induces LTP in anesthetized adult rats, but the mechanisms of LTP maintenance remain unclear^[34]. In this study, we used HFS to further understand the underlying mechanisms of the LTP enhancement of PGS32. The intensity of LTP after PGS32 injection was significantly enhanced compared to the control group after HFS. These results suggest that PGS32 not only can induce LTP but also can maintain LTP. PGS32 may promote synaptic plasticity in the hippocampus. Although we demonstrated that PGS32 has the ability to improve learning and memory and to promote hippocampal LTP, further studies in other animal models of dementia are necessary to confirm the protective effects of PGS32.

In summary, we have validated that PGS32, a triterpene saponin from Polygalae Radix, can ameliorate memory impairments induced by scopolamine in mice. The mechanisms by which PGS32 improves memory dysfunction appear

to involve the protection of neurons from glutamate and ROS damage, the up-regulation of the phosphorylation of TrkB and the maintenance of LTP. These results suggest that PGS32, an active constituent of *Polygalae Radix*, may be a competitive candidate for the treatment of neurodegenerative disorders.

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Author contribution

Heng ZHOU analyzed the data and wrote the paper; Wei XUE and Shi-feng CHU performed the research; Zhen-zhen WANG provided assistance with experimental methods and revised the paper; Dong-ming ZHANG and Chuang-jun LI extracted the PGS32; Yi-na JIANG, Lin-ming LUO and Piao LUO took part in the research; Gang LI provided comments and technical support; Nai-hong CHEN designed the study and revised the paper.

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