The effect and mechanism of combination of total paeony glycosides and total ligustici phenolic acids against focal cerebral ischemia

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The root of Paeonia lactiflora Pall. (Chishao, CS) and Ligusticum chuanxiong Hort. (Chuanxiong, CX) were widely used as a drug pair in Chinese Medicine, and the combination of CS and CX showed a more significant inhibition on neuronal apoptosis in our previous study. In the present study, total paeony glycosides (TPGs) from CS and total ligustici phenolic acids (TLPAs) from CX were combined to evaluate the synergistic effects against focal cerebral ischemia both in vitro and in vivo. The combination of TPGs and TLPAs at 7:3 had the best anti-oxidative stress and anti-inflammatory effect on OGD-induced HUVEC. Additionally, the infarction area proportion and neuron apoptosis of rats by TPGs:TLPAs (7:3) was significantly lower than their alone in MCAO rats. Moreover, TPGs:TLPAs of 7:3 showed a more significant effect on decreasing the expression of MMP-2 and MMP-9, and increasing the protein expression or mRNA level of TIMP-1 than other combinations. The optimal ratio of TPGs and TLPAs at 7:3 could bring more remarkable protective effects against focal cerebral ischemia in MCAO rats by alleviating oxidative stress, inflammatory and neuronal apoptosis to protect the blood-brain barrier. Overall, the present study provided beneficial evidence for clinical application of CS and CX as a “drug pair”.

As the third highest worldwide cause of mortality1, stroke has been treated using antithrombotic therapy2, anticoagulant therapy3,4, neuroprotective and vascular dilated therapy. Numerous studies have demonstrated that the severity of neuronal apoptosis was associated with neurofunction, indicating a crucial role in the prognosis of ischemic stroke sufferer1. Early anti-inflammatory and anti-oxidative stress may promote apoptosis of nerve cells. Thus attenuating these potential damage of nerve cells in ischemic area will be an effective strategy in clinical application.

Chinese medicine (CM) has been used to treat stroke for a long time. Many classical prescriptions for treating cerebral diseases, such as Xiao-xu-ming decoction6 are still being commonly used in modern clinical both in China and in other countries8. Paeonia lactiflora Pall. (Chinese name: Chishao, CS) has been used worldwide for centuries and reported to exhibit various pharmacological bioactivities including anticoagulation, cognition enhancement, immunoregulatory9 and anticance10, antioxidative11, and antidiabetic properties12. The ischemic stroke protection of Paeoniae Radix may be related to its anti-oxidative stress and improvement of brain micro-circulation. Ligusticum chuanxiong Hort. (Chinese name: Chuanxiong, CX) which belongs to the Umbelliferae family, is a valuable medicinal herb used worldwide and has been used for the treatment of atherosclerosis, hypertension, angina pectoris, and anti-inflammatory via acilitating blood circulation and dispersing blood stasis13. Our previous study revealed the combination of CS and CX exerted more remarkable protective effects than alone by attenuating ER-stress-dependent apoptotic signaling and protected the blood-brain barrier (BBB)14.

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Glycosides are the main active constituents in CS, while phenolic acids are the major active components in CX. However, the best efficacy of these components might be associated with its combination at the optimal ratio. In current study, we use the ischemic model both in vitro and in vivo to filter the best combination of total paony glycosides (TPGs) and total ligustici phenolic acids (TLPAs) against focal cerebral ischemia.

### Results

#### Cell viability.

MTT assay was used to evaluate cell viability for screening the optimal drug concentration of TPGs and TLPAs for Human Umbilical Vein Endothelial Cells (HUVEC) cells. As shown in Table 1, the viability of HUVEC cells ranged from 53.72 ± 2.99% to 101.61 ± 2.11% when treated with different concentration of TPGs from 1 × 10⁻³ g/mL to 1 × 10⁻⁵ g/mL. While the viability of HUVEC cells ranged from 72.37 ± 5.01% to 99.96 ± 3.25% when treated with TLPAs from 1 × 10⁻⁴ g/mL to 1 × 10⁻⁵ g/mL. Thus 2 × 10⁻⁴ g/mL TPGs with the cell viability of 96.79 ± 1.98% and 2 × 10⁻⁴ g/mL TLPAs with cell viability of 95.01 ± 2.98% were chosen for further experiments.

#### Anti-oxidative stress activity of TPGs:TLPAs in MCAO rats.

As shown in Fig. 1A, infarction areas were significantly increased. (P < 0.01, Fig. 1B). When TPGs:TLPAs at the best optimal ratio on MCAO rats. Improvement of TPGs:TLPAs at 7:3 on neurological deficit score of MCAO rats. As the results showed in Table 4, at 24 h after cerebral ischemia, the neurological deficit score of the sham-operated rats was 0, indicating that the sham-operated rats did not have neurological impairment symptoms. Compared with the sham-operated group, the neurological deficit scores in their combination of different ratio. A more significant decrease of these inflammatory cytokines was observed in HUVEC treated with TPGs and TLPAs combination at the ratio of 7:3 (P < 0.01, P < 0.05).

### Anti-inflammation of different ratio with TPGs and TLPAs in OGD-induced HUVEC.

As depicted in Table 3, the contents of TNF-α, IL-6 and sICAM-1 were significantly increased after modeling (P < 0.01, P < 0.05). The contents of TNF-α, IL-6 and sICAM-1 were all decreased in different degrees after treatment of TPGs, TLPAs and their combination of different ratio. A more significant decrease of these inflammatory cytokines was observed in HUVEC treated with TPGs and TLPAs combination at the ratio of 7:3 (P < 0.01, P < 0.05).

### The effect of TPGs and TLPAs at the best optimal ratio on MCAO rats.

Improvement of TPGs:TLPAs at 7:3 on infarction area of MCAO rats. As shown in Fig. 1A, infarction areas were shown in white. There was no infarction in the sham operation group, while infarction occurred in each group after the model was established. The cerebral infarct size ratio of the model group was reached 49.72%, which was significantly different from the sham operation group (P < 0.01). Compared with the model group, the infarct size of cerebral ischemia rats was significantly reduced after administration (P < 0.01, Fig. 1B). When TPGs:TLPAs at the ratio of 7:3, the treatment of infant size in cerebral ischemia rats was most effective, showed a significantly lower in infarction areas than other combinations (P < 0.01, P < 0.05). There was no difference in cerebral infarct size ratio between TPGs:TLPAs treatment at 5:5 and TPG treatment alone, but the effect was better than that of the TLPAs alone group (P < 0.05).

Anti-oxidative stress activity of TPGs:TLPAs at 7:3 in MCAO rats. Anti-oxidative stress activity of TPGs:TLPAs combinations in MCAO rats was presented in Table 5. Comparing with sham control group, the activities of SOD, CAT and GSH-Px of model rats were significantly decreased (P < 0.01, P < 0.05), while the levels of MDA, and LPO were significantly increased (P < 0.01, P < 0.05). The activities of SOD, CAT, GSH-Px, MDA, and LPO were significantly increased (P < 0.01). Compared with the model group, the infarct size of cerebral ischemia rats was significantly reduced after administration (P < 0.01, Fig. 1B). When TPGs:TLPAs at the ratio of 7:3, the treatment of infant size in cerebral ischemia rats was most effective, showed a significantly lower in infarction areas than other combinations (P < 0.01, P < 0.05). There was no difference in cerebral infarct size ratio between TPGs:TLPAs treatment at 5:5 and TPG treatment alone, but the effect was better than that of the TLPAs alone group (P < 0.05).

### Table 1. Effect of TPG and TLPA at different concentrations on the survival rates of HUVEC (x ± SD, n = 6).

| Group   | Dosage (g/mL) | OD       | Cell viability (%) |
|---------|---------------|----------|--------------------|
| Control blank | —            | 0.4328 ± 0.1435 | 100                |
| TPGs    | 10⁻³          | 0.2325 ± 0.0247 | 53.72 ± 2.99       |
|         | 5 × 10⁻⁴      | 0.2765 ± 0.0427 | 63.88 ± 2.64       |
|         | 2 × 10⁻⁴      | 0.4189 ± 0.0167 | 96.79 ± 1.98       |
|         | 10⁻³          | 0.4203 ± 0.0379 | 97.12 ± 3.18       |
| TLPAs   | 10⁻³          | 0.4398 ± 0.0622 | 101.61 ± 2.11      |
|         | 10⁻²          | 1.0437 ± 0.1038 | 105.11 ± 3.11      |
|         | 10⁻¹          | 2.0437 ± 0.2038 | 110.11 ± 5.11      |

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Table 2. Effect of TPGs and TLPAs on T-SOD, T-NOS, i-NOS, CAT, LPO, MPO and GSH-PX activity and MDA, NO content in OGD-induced HUVEC (± SD, n = 3). Notice: *P < 0.01 or **P < 0.05, compared with control group; *P < 0.01 or **P < 0.05, compared with model group.

| Group | TPGs | TLPAs | TPGs:TLPAs | T-SOD (U/mL) | T-NOS (U/mL) | i-NOS (U/mL) | CAT (U/mL) | LPO (μmol/L) | MPO (U/L) | GSH-PX (μmol/L) |
|-------|------|-------|-------------|---------------|---------------|--------------|-------------|--------------|-------------|----------------|
| Control | 8.67 ± 0.12 | 5.77 ± 0.07 | 3:2 | 2182.19 ± 1.19 | 1199.13 ± 0.07 | 525.09 ± 0.12 | 12.26 | ± 0.03 | 2.87 | ± 0.03 |
| Model | 4.13 ± 0.17 | 3.50 ± 0.07 | 9:1 | 2182.19 ± 1.19 | 1199.13 ± 0.07 | 525.09 ± 0.12 | 12.26 | ± 0.03 | 2.87 | ± 0.03 |
| Ni | 3.79 ± 0.24 | 3.50 ± 0.07 | 8:2 | 2182.19 ± 1.19 | 1199.13 ± 0.07 | 525.09 ± 0.12 | 12.26 | ± 0.03 | 2.87 | ± 0.03 |

Table 3. Effect of XS on TNF-α, IL-6, sICAM-1 expression in OGD-induced HUVEC (± SD, n = 3). Notice: *P < 0.01 or **P < 0.05, compared with control group; *P < 0.01 or **P < 0.05, compared with model group.

| Groups | 24 h | 1 week | 2 week |
|--------|------|--------|--------|
| Sham | 0 | 0 | 0 |
| Model | 3.32 ± 0.09** | 3.35 ± 0.13** | 3.34 ± 0.13** |
| Ni | 3.30 ± 0.17** | 2.23 ± 0.21** | 2.08 ± 0.17** |
| TPGs | 3.29 ± 0.22** | 2.62 ± 0.20** | 2.30 ± 0.20** |
| TPGs:TLPAs (7:3) | 3.30 ± 0.18** | 2.48 ± 0.13** | 2.21 ± 0.22** |
| TPGs:TLPAs (5:5) | 3.28 ± 0.23** | 2.61 ± 0.06** | 2.29 ± 0.19** |
| TLPAs | 3.31 ± 0.19** | 2.87 ± 0.13** | 2.69 ± 0.11** |

Table 4. The neurological function of rats after MCAO (± SD, n = 24). Notice: *P < 0.01, compared with sham group; *P < 0.01 or **P < 0.05, compared with model group; *P < 0.01 or **P < 0.05, compared with TLPAs group; *P < 0.05, compared with TPGs group; %P < 0.05, compared with TPGs:TLPAs (5:5) group; @P < 0.01 or $P < 0.05, compared with the previous group.

were significantly reversed by the treatment with Ni, TPGs, TLPAs, TPGs:TLPAs at 7:3 and TPGs:TLPAs at 5:5. Importantly, TPGs:TLPAs at 7:3 ratio has a more significant amelioration on these activities or levels, compared with other combinations (P < 0.01, P < 0.05).
Effects of TPGs:TLPAs at 7:3 on functional recovery and anti-apoptosis of MCAO rats. Clear and integral tissue structure was observed in sham group with clear neuronal nuclei, relatively intact nuclear membrane and normal glial cells and capillary morphogenesis (Fig. 2C). However, in model group, edema of the neuropile were observed in model group, and most neurons in the damage areas appeared shrunken with eosinophilic cytoplasm and triangulated pyknotic nuclei. Histopathological abnormalities were significantly improved by TPGs and TLPAs combinations. Particularly, the treatment of TPGs:TLPAs (7:3) had the most obvious effect on improving the ischemic injury compared with other groups ($P < 0.01, P < 0.05$).

In sham control group, polygons neurons, abundant cytoplasm, clear nucleoli and integral cell membrane were observed by TEM (Fig. 1D) by TEM. After MCAO modeling, neurons were swelled, organelles were broken...
or disappeared, and cell membranes were adhered with synapses. Meanwhile myelin in most synapses from surrounding stroma was disappeared (red arrow) or swelled (yellow arrow). After being treated with Ni, TPGs, TLPAs, TPGs:TLPAs at 7:3 and 5:5, apoptotic neurons were significantly ameliorated. Especially, the treatment with TPGs:TLPAs (7:3) had the most obvious improvement on neurons apoptosis compared with other combinations ($P < 0.01$, $P < 0.05$).

### Protection of TPGs:TLPAs at 7:3 on blood-brain barrier in MCAO rats

As shown in Table 6, comparing with sham control group, the activities of MMP-9 and PAI-1 were significantly decreased in model group ($P < 0.01$), whereas the activities of PA were significantly increased ($P < 0.01$). After intervening by Ni, TPGs, TLPAs, TPGs:TLPAs at 7:3 and 5:5, the activities of MMP-9 and PAI-1 were significantly increased, whereas the activities of PA was significantly decreased. TPGs and TLPAs at the ratio of 7:3 significantly changed the activity to a higher level when compared with other combinations ($P < 0.01$, $P < 0.05$).

### Protection of TPGs:TLPAs at 7:3 on apoptotic protein in MCAO rats

After modeling, protein expression of Bax and caspase-3 were significantly enhanced whereas Bcl-2 was apparently declined ($P < 0.01$). Compared with model group, protein expressions of Bax and caspase-3 in rats brain decreased significantly while Bcl-2 was significantly increased after treatment the with Ni, TPGs, TLPAs, TPGs:TLPAs at 7:3 and 5:5 for two weeks ($P < 0.01$, $P < 0.05$) (Fig. 3A). It was important that protection of TPGsTLPAs at 7:3 on apoptosis in MCAO rats was significantly better than any other groups ($P < 0.01$, $P < 0.05$).

### Discussion

TCM, consisting of multiple herbs, was beneficial to synergistic effect of multi-components for the treatment and prevention of diseases. In traditional application, CS (Paeonia lactiflora Pall.) and CX (Ligusticum chuanxiong Hort.) were generally used as “drug pair” to treat the stasis and stroke. As known, CS was used for “eliminating blood stasis and relieving pain” (Chinese Pharmacopoeia 2015). Modern phytochemical and pharmacological studies have shown that monoterpenic glycosides, galloyl glucoses and phenolic compounds are the major bioactive components of CS. Research showed its major compound paenoflorin was beneficial for anti-inflammatory and autoimmune disease treatment. In this study, organic acids including ferulic acid and caffeic acid were isolated and identified from Ligusticum Chuanxiong Hort. These components have been proved to hold chemoprevention on apoptosis and reduction on the risk of ROS-mediated diseases. According to our previously study, the combination of CS and CX have potential effects on curing cerebral disease or stroke. However, the best efficacy of these components might be associated with its combination with the optimal ratio. Total paeony glycosides (TPGs) are the main active constituents in CS and total ligustici phenolic acids (TLPAs) are the major active components in CX. TPGs were reported to have a good effect on both cerebral ischemia and cardiac ischemia.

| Groups       | SOD (U/mL) | CAT (U/mL) | GSH (μmol/L) | MDA (U/mL) | LPO (μmol/L) |
|--------------|------------|------------|--------------|-------------|--------------|
| Sham         | 198.68 ± 2.26 | 12.98 ± 1.01 | 410.23 ± 19.17 | 38.12 ± 8.71 | 5.09 ± 0.73 |
| Model        | 121.33 ± 5.40 | 9.01 ± 1.15 | 391.56 ± 14.69 | 19.38 ± 3.69 | 10.12 ± 1.29 |
| Ni           | 191.36 ± 14.30 | 10.11 ± 1.82 | 397.16 ± 14.33 | 41.11 ± 8.76 | 5.28 ± 0.38 |
| TPGs         | 179.35 ± 4.97 | 8.31 ± 0.87 | 389.67 ± 10.91 | 50.69 ± 12.65 | 6.82 ± 1.75 |
| TPGs:TLPAs (7:3) | 189.56 ± 2.66 | 10.12 ± 1.29 | 395.37 ± 19.38 | 43.77 ± 7.41 | 0.55 ± 0.68 |
| TPGs:TLPAs (5:5) | 180.72 ± 3.69 | 9.01 ± 1.15 | 391.56 ± 14.69 | 19.38 ± 3.69 | 10.12 ± 1.29 |
| TLPAs        | 173.18 ± 14.30 | 8.39 ± 1.29 | 389.15 ± 19.38 | 50.36 ± 7.81 | 8.05 ± 0.68 |

Table 5. The effect of XS on SOD, CAT, GSH-Px activity and LOP, MDA content in rats serum of different treatment groups after cerebral ischemia (X ± SD, n = 24). Notice: **P < 0.01 or *P < 0.05, compared with sham group; ***P < 0.01 or **P < 0.05, compared with model group; &P < 0.01 or &P < 0.05, compared with TLPAs group; ##P < 0.01 or ##P < 0.05, compared with TPGs group; ##P < 0.01 or &P < 0.05, compared with TPGs:TLPAs (5:5) group.
In current study, we use the ischemic model both in vitro and in vivo to identify the best combined ratio of TPGs and TLPAs against focal cerebral ischemia. The optimal ratio of TPGs and TLPAs at 7:3 could bring more remarkable protective effects against focal cerebral ischemia in MCAO rats by alleviating oxidative stress, inflammatory and neuronal apoptosis to protect the blood-brain barrier. The present study provided beneficial evidence for clinical application of CS and CX as a "drug pair".

Cerebral disease or stroke is caused by lack of oxygen to the cerebrum, however their pathophysiology are still not completely clear. Cerebral ischemia, with risk factors including hypertension, diabetes, hyperlipidemia,
smoking and tiredness, may cause irreversible structural changes in a specific vascular territory. However, the effect of cerebral ischemia on BBB, a diffusion barrier essential for maintenance of homeostasis and normal function of the central nervous system, has been broadly investigated. In current study, we investigated the protection of the combination of TPG and TLPA on BBB by anti-inflammation, anti-oxidative stress and anti-apoptosis effects.

Table 6. MMP-9, PA, PAI-1 content in different treatment rats serum after MCAO (x ± SD, n = 24). Notice: \( \text{"P} < 0.01 \) or \( \text{"P} < 0.05 \), compared with sham group; \( \text{"P} < 0.01 \) or \( \text{"P} < 0.05 \), compared with model group; \( \text{"P} < 0.01 \) or \( \text{"P} < 0.05 \), compared with TLPA group; \( \text{"P} < 0.01 \) or \( \text{"P} < 0.05 \), compared with TPGs group; \( \text{"P} < 0.01 \) or \( \text{"P} < 0.05 \), compared with TPGs:TLPA (5:5) group.

| Groups          | MMP-9 (ng/mL) | PA (mg/mL) | PAI-1 (ng/mL) |
|-----------------|---------------|------------|---------------|
| Sham            | 38.12 ± 1.87  | 241.91 ± 26.19 | 51.13 ± 9.71  |
| Model           | 60.03 ± 1.91**| 67.36 ± 18.01**| 185.83 ± 10.12**|
| Ni              | 42.05 ± 1.33**| 221.46 ± 15.43**| 69.27 ± 9.31**|
| TPGs            | 50.43 ± 1.51* | 191.52 ± 16.51**| 107.42 ± 10.19* |
| TPGs:TLPA (7:3) | 45.33 ± 2.01** | 219.93 ± 19.06** | 95.37 ± 8.76** |
| TPGs:TLPA (5:5) | 48.62 ± 1.88** | 201.83 ± 20.31** | 103.13 ± 13.51** |
| TLPAs           | 50.43 ± 1.92**| 190.72 ± 17.31** | 110.19 ± 10.03** |

Figure 3. Effect of TPGs and TLPA on Bax, Bcl-2 and caspase-3 protein expression in cerebral ischemia rats. (A) Images of western blot; (B) Relative protein expression of Bax, Bcl-2 and caspase-3; (C) The mRNA level in cerebral ischemia rats. \( \text{"P} < 0.01 \), compared with sham group; \( \text{"P} < 0.01 \) or \( \text{"P} < 0.05 \), compared with model group; \( \text{"P} < 0.01 \) or \( \text{"P} < 0.05 \), compared with TLPA group; \( \text{"P} < 0.01 \) or \( \text{"P} < 0.05 \), compared with TPGs group; \( \text{"P} < 0.01 \) or \( \text{"P} < 0.05 \), compared with TPGs:TLPA (5:5) group.
Because of complicated factors for cerebral disease or stroke, it is inevitably to be a comprehensive pathological process. Cerebral inflammation participated in all stages of ischemic cascade which plays a crucial role in the pathophysiology of ischemic stroke, while oxidative stress has also been reported to be one of the main cause in ischemia reperfusion injury.

Matrix metalloproteinases (MMPs), a family of zinc-binding proteolytic enzymes, contribute to the pathology of cerebral infarction by degrading a number of extracellular matrix molecules to breakdown the blood-brain barrier. MMPs-mediated BBB disruption has been found to be a critical pathological mechanism of cerebral disease or stroke. However, this activation of MMPs is complexly modulated by transcriptional regulators and inhibitor tissue inhibitor of metalloproteinases (TIMPs). Under normal physiological conditions, MMPs are directly responsible for pathogenesis of cerebrovascular diseases and their activity is regulated by TIMPs. The evidence showed that MMP-2 and MMP-9 might be up-regulated and activated during cerebral disease or stroke which are expressed particularly frequently in the nervous system. MMP-2 also plays a significant role in the regulation of activity of platelets, while MMP-9 damages matrix components of the basement membrane leading to neuro-inflammation after focal cerebral ischemia by maintaining the integrity of cerebral vasculature. MMP-9, one of the most important MMPs correlated with BBB damage, accompanied with its endogenous inhibitor TIMP-1, has been confirmed to be important in vascular remodeling and neuro-protective effect. This study indicated that there was a more significant effect of TPGs:TLPAs (7:3) on adjusting MMP-2, MMP-9 and TIMP-1 protein expression and mRNA level than other groups.

It has been proved that apoptosis is associated with cerebral disease or stroke leading to BBB dysfunction, inflammation and oxidative stress resulting in brain damage. Histopathological changes and TEM images in current study indicated that TPGs and TLPAs especially the TPGs:TLPAs at 7:3 could alleviate neuronal apoptosis.

Conclusions
In this study, the optimal ratio of TPGs and TLPAs on improving cerebral ischemia damage was compared and screened according to ODG-induced HUVEC cells in vitro and MCAO rats in vivo. The results demonstrated the combination TPGs and TLPAs at 7:3 might be more beneficial to this protective effect via alleviating oxidative stress, inflammation and neuronal apoptosis. This study provided an evidence of the combination of CS and CX for clinical application as “drug pair”, and more importantly, the study revealed that the proper ratio of TPGs and TLPAs could be critical for CS-CX “drug pair” in curing focal cerebral ischemia.

Materials and Methods
Materials and chemicals. *Paeonia lactiflora* Pall. (CS) was offered by Sichuan new Lotus Chinese Herbal Medicine Co., Ltd. (Chengdu, China) with the lot number 1311060. *Ligusticum chuanxiong* Hort. (CX) was got from Songling Chinese Herbal Medicine Co., Ltd. (Nanjing, China) with the lot number 131017. The pharmaceutical botanies of the medicinal material were identified by Prof. Dekang Wu from Nanjing University of Chinese Medicine (Nanjing, China).

Nimodipine (Ni, approval number was H20003010), as a positive control drug, was obtained from Bayer Healthcare (Beijing, China). Kits for SOD, MDA, CAT, GSH-Px and LPO assay were purchased from Jiancheng Bioengineering Institute (Nanjing, China). ELISA kits for MMP-9, PA and PAI-1 were provided by Beyotime Biotech. Co., LTD. (Shanghai, China). Hematoxylin-eosin staining solution was offered by Shanghai Yuanmu Biotechnology Co., Ltd. (Shanghai, China). RPMI 1640 medium and trypsin were provided by KeyGEN Biotech Co., Ltd (Nanjing, China). Fetal bovine serum (FBS) was purchased from Gibco/BRL (Grand Island, USA).
5-dimethyl-2-thiazolyl)-2, 5-diphenyl-2-H-tetrazolium bromide (MTT) and dimethyl sulfoxide (DMSO) were provided by Sigma (St. Louis, USA). Rabbit anti-mouse Bax, Bcl-2, and TIMP-1 monoclonal antibodies were offered by Santa Cruz Biotechnology, Inc. (Dallas, USA). Rabbit anti-mouse MMP-9, PERK, XBP-1, ATF-6 and CHOP monoclonal antibodies were obtained from Abcam, Inc. (Cambridge, UK). Other rabbit anti-mouse monoclonal antibodies were got from Boster Biological Engineering Co., Ltd. (Wuhan, China). EliVision plus and DAB kits were offered by MXB Biological Technology Co., Ltd. (Fuzhou, China).

Preparation of TPGs and TLPAs. The preparation of TPGs was performed as previous studies with modifications. Pieces of CS with 1 kg were weighed and extracted twice (1.5 h/time) with 10 times amount of 70% (v/v) ethanol in reflux extraction device. The extraction was concentrated to 1 g decocion pieces/mL. TPGs was defined as the 30% ethanol eluant chromatographed with a D-101 macroreticular resin (Cangzhou Bon Adsorber Technology Co., Ltd, Cangzhou, China). On the basis of previous studies32, 1 kg CX were weighed and extracted twice (1.5 h/time) with 15 times amount of 80% (v/v) ethanol in reflux extraction device. The extraction was concentrated to 1 g decocion pieces/mL and was chromatographed with AB-8 macroreticular resin (Cangzhou Bon Adsorber Technology Co., Ltd, Cangzhou, China) column. Finally, the eluant of 30% ethanol was defined as TLPAs. The yield of CS and CX extraction was 17.81% and 16.63%, respectively. The content of TPGs was 3.62% after extraction with 70% ethanol. And after purification by macroporous resin, the content of the TPGs used in the study was 47.92% finally. While the content of TLPAs was 2.42% after extraction with 80% ethanol, and the content of the TLPAs used in the study was 41.43% finally after purification (The data shows in Supplementary Information).

Cells culture. HUVEC was purchased from Shanghai Institute of Biochemistry and Cell Biology (Shanghai, China) and cultured in an incubator with 5% CO2 at 37°C. Cells were treated with RPMI 1640 medium containing 80 units/mL of penicillin, 0.08 mg/mL of streptomycin (KeyGEN, China) and 10% fetal bovine serum (FBS, Gibico, USA).

Cell proliferation assay by MTT. After 90% confluence, cells were digested by 0.25% trypsin-0.02% EDTA for the passage. A cell density of 5000 cells/well was seeded into 96-well plates (100 μL). Cells were treated with different concentrations of TPGs prepared (10−3, 5 × 10−3, 2 × 10−4 and 10−5 g/mL) and TLPAs (10−3, 5 × 10−4, 2 × 10−4 and 10−5 g/mL) for 24 h respectively. MTT stock solution of 5.0 mg/mL (10 μL) was added to each well for 4 h to form water-insoluble purple crystal formazan. At the end of incubation, DMSO of 150 μL was added for 10 min microvibration after removing medium. The absorbance was measured at 570 nm on a microplate reader (Thermo, New York, USA).

Cell OGD model and drug treatment. HUVECs modeled with oxygen glucose deprivation (OGD) was used as an ischemic model in vitro according to previous reports. Cells of 90% confluence were cultured in 96-well plates (5000 cells/well, 100 μL). For simulated ischemia-reperfusion, cells were placed in a hypoxic incubator (5% CO2, 95% N2) at 37°C for 4 h.

Then serum-free RPMI 1640 medium was used in control group and model group. The other OGD cells were divided into 12 groups (n = 6/group) including positive control group (Nimodipine 10 μM); TPGs group; TLPAs group; TPGs:TLPAs at 9:1, 8:2, 7:3, 6:4, 5:5, 4:6, 3:7, 2:8 and 1:9 ratios. The treatment dosage was 2 × 10−4 g/mL.

Animal model and drug treatment. Male Sprague-Dawley rats (280 ± 20 g, licence number: SCXK (Su) 2015–0001) were supplied by Nantong University (Nantong, China). Rats were kept at a temperature of 25°C and the relative humidity of 45%. All the experiment procedures were approved by the Institutional Animal Care and Use Committee of the Jiangsu Provincial Academy of Chinese Medicine in accordance with published National Institutes of Health guidelines, the ethics code number of our animal experimental was ZYY20151207. Before modeling, rats were housed for 7 d to adapt to the environment on a 12 h/12 h light/dark cycle with food and water ad libitum.

The middle cerebral artery occlusion (MCAO) model was performed as described previously. Rats were injected intraperitoneally 10% chloral hydrate (300 mg/kg) for anesthetizing. After anesthetized, rats were laid in dorsal recumbency and the right side of carotid artery was exposed and isolated. The origin of the right middle cerebral artery (MCA) was occluded by inserting a 0.26 mm heparin-dampened monofilament nylon suture (Ethicon, Inc., Osaka, Japan) with a heat-rounded tip from the external carotid artery (ECA) to the right internal carotid artery (ICA). Thread at the distal end of common carotid artery (CCA) was fastened when the suture was advanced closed to the origin of the MCA. Extravascular suture was cut before the closed neck incision. After preventing wound infection, rats in the sham group were performed with the same surgical procedures without insertion of the heparin–dampened monofilament nylon suture. The body temperature of these rats was maintained at 37°C during surgery. After surgery, sodium penicillin (100,000 U/rat/d) was given to prevent infection for three days successively.

Neurological examination was conducted to screen the successful models after modeling for 24 h. The successfully modeled rats were randomly divided into 7 groups (n = 24/group): model group (MCAO, Saline 10 mL/kg); sham control group (Saline 10 mL/kg); positive control group (Nimodipine 7.53 mg/kg); TPGs group; TLPAs group; TPGs:TLPAs at 7:3 group and TPGs:TLPAs at 5:5 group. The treatment drug dose was 500 mg/kg according to the previous research17. Equivalent saline was given to the sham rats and model rats. All rats were treated by gavage administration for two weeks consecutively.
Behavioral measurements. Neurological evaluation was performed by assessing the neurological deficit score \(^{3,4} \) 24 h after modeling and every week, with each point characterized by the following attributes: 0 = no symptoms of neurological dysfunction; 1 = failure to extend left forelimb fully; 2 = circling to the contralateral side; 3 = falling to the left; 4 = no spontaneous walk or in a comatose state; 5 = death.

Infarct volume and area measurement. Rats were sacrificed by deeply anaesthetized with 5% isoflurane and euthanized by cervical dislocation. Brain tissues were kept at \( -20 \) °C for 30 min after ice cold saline washing. Coronal sections (3 mm) of the frozen brains were cut with the sharp blades and stained with 2% 2, 3, 5-triphenyltetrazolium chloride (TTC, dissolved in PBS solution, pH 7.4) at 37 °C for 30 min in the dark. Sections were reversed every 5 min to keep uniform coloring of the sections. Viable tissues were stained in deep red while the infarcts remain unstained. TTC-stained sections were photographed with a digital camera after being fixed with 4% paraformaldehyde. Finally, images were captured by a Nikon D700 digital camera (Nikon, Japan).

The infarct volume of each section was calculated using Image-Pro Plus 6.0 image analysis system (Media Cybernetics, USA). The infarct area (IS) was calculated as previously described using the following equation \(^{35} \):

\[
\text{IS} = (1 - S_1/S_2) \times 100\% \quad (S_1: \text{ipsilateral hemisphere non-infarcted area}; S_2: \text{contralateral hemisphere area})
\]

Hematoxylin-Eosin (HE) staining. After rats being sacrificed by isoflurane, the brains of rats were taken out and fixed in 4% (w/v) paraformaldehyde for 3 days. The brain block was embedded in paraffin and cut into 3 mm coronal sections. Sequentially, the sections were stained with hematoxylin and eosin (HE), dehydrated, vitrified and mounted. The program of staining was observed under the IX51 microscope (Olympus Corporation, Japan).

Content measurement of SOD, CAT, GSH-Px, LPO and MDA. Cells or serum of rats was collected for the content determination according to the manufacturer’s protocols. The optical density (OD) of samples was obtained with microplate reader (Thermo Scientific, MA, USA) at wavelength of 550 nm (SOD), 405 nm (CAT), 412 nm (GSH-Px), 586 nm (LPO) and 532 nm (MDA) respectively.

Enzyme-linked immunosorbent assay (ELISA) for MMP-9, PA and PAI-1. The levels of MMP-9, PA and PAI-1 in serum of MCAO rats were determined with ELISA according to manufacturer’s instructions. The samples were readed at 450 nm using a microplate reader (Thermo Scientific, MA, USA). The contents of MMP-9, PA and PAI-1 were calculated according to the standard curve.

Immunohistochemistry. Brain tissues were removed and immediately immersed in formaldehyde solution. Paraffin embedded tissues were sectioned (5 μm thick) and labeled with 3% H\textsubscript{2}O\textsubscript{2}, 3% normal goat serum. Briefly, sections were incubated with anti-mouse antibodies overnight. Then secondary antibodies, secondary biotinylated conjugates and diaminobenzidine were added as the instructions of DAB kit. Finally, slides were dehydrated, cleared, and mounted for visualization using IX51 microscope (Olympus Corporation, Japan).

Transmission electron microscope (TEM). Preparation of samples was performed according to previous method. Sections of 1 mm\textsuperscript{3} were taken out from hippocampal CA1 region of rats’ brain and fixed with 5% glutaraldehyde followed by PBS (pH = 7.4) washing. Then sections were postfixed in 1% osmic acid and dyed with 2% uranyl acetate. Sequentially, sections were dehydrated by 50%, 70%, 90% and 100% acetone and embedded. Samples were analyzed using the JEM-1200EX transmission electron microscope (JEOL, Japan) at a voltage of 80kV.

Western blotting analysis. Protein levels in brain tissues were determined by Western blotting analysis. Briefly, proteins were extracted from tissue homogenates and an equal amount of total protein was separated by 10% sodium dodecy1 sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and then transferred from the SDS-PAGE gel to PVDF membrane (Millipore, USA). After being blocked with 5% BSA in tris-buffered saline Tween-20 (TBST), membranes were incubated with the primary antibodies at a dilution of 1:1000 overnight. Subsequently, membrane was washed (three times, 5 min), and incubated with secondary antibody (1:5000) at 37 °C for 30 min. The blots were visualized with ECL-Plus reagent (Santa Cruz, USA) and analyzed with Image pro plus (IPP 6.0, Media Cybernetics, MD, USA) software. β-actin was used as loading control.

Quantitative PCR (Q-PCR). Total RNA of all brain samples was extracted by TRIzol reagent. Then it was reverse transcribed with a SuperScript III First-Strand Synthesis System for quantitative real-time polymerase chain reaction (q-PCR) following manufacturer’s indications. The Primer sequences used in this study were shown in Table 7. SYBR green of 10 μL, 6 μL molecular grade water, 1 μL of each forward and reverse primers and 2 μL cDNA were contained in each reaction of the PCR plate. The amplification was performed under the following conditions: 10 minutes at 95°C, 50 cycles at 95°C for 15 seconds and 60°C for 60 seconds. Q-PCR was performed under standard conditions and all experiments were run in triplicate. GAPDH was used as the internal reference.

TUNEL staining. Cell apoptosis in the brain was detected by TUNEL staining according to the manufacturer’s protocol (KeyGEN, Nanjing, China). Briefly, 5 μm slide sections were deparaffinized in Histoclear, rehydrated and treated with NaCl (0.85% w/v) for 5 min, then washed with 4% w/v paraformaldehyde and permeabilized with Proteinase K. After that sections were equilibrated and incubated with Biotinylated Nucleotide mix and recombinant Terminal Deoxynucleotidyl Transferase (rTdT) at 37 °C for 1 h till the
reaction was stopped. Then endogenous peroxidases blocked with H2O2. Sections were incubated with Streptavidin–horseradish peroxidase conjugate solution and colored with a diaminobenzidine substrate. Sections were processed in parallel without rTdT as controls. Each section was visualized with light microscopy and images captured digitally and analyzed.

**Statistical analysis.** All data were expressed as means ± standard deviation (SD). Statistical analysis was performed by SPSS 16.0 with one-way analysis of variance (ANOVA) based on Student’s two-tailed unpaired t-test or Dunnett’s multiple comparisons test. The value of P less than 0.05 was considered to be a statistically significant difference. Immunohistochemistry images and WB images were analyzed by Image-Pro Plus 6.0 (Media Cybernetics, MD, USA), a software specializes in image processing.

**Chemical compounds studied in this article.** Protocatechuic (PubChem CID: 72); Oxypaeoniflorin (PubChem CID: 46882883); Catechin (PubChem CID: 73160); Albiflorin (PubChem CID: 51346141); Caffeic acid (PubChem CID: 689043); Paeoniflorin (PubChem CID: 75124163); Ferulic acid (PubChem CID: 445858); Benzoyl paeoniflorin (PubChem CID: 73157327).

Received: 12 July 2019; Accepted: 5 February 2020; Published online: 28 February 2020

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Acknowledgements
All authors gratefully acknowledge the National Natural Science Foundation of China (81573620, 81603382, 81973536, 81904059). Natural Science Research in Jiangsu Province(19KBJ360015, BK20190803). A Project Funded by the Priority Academic Program Development of Jiangsu Higher Education Institutions(Integration of Chinese and Western Medicine). Program for Innovative Research Team of Six Talent Peaks Project in Jiangsu Province (SWYY-CXTD-004), Scientific Research Project of State Traditional Chinese Medicine Clinical Base Construction (JDZX2015072), Jiangsu Postdoctoral Fund (1601023 C), China Postdoctoral Fund (2016M601864), and “Double First-Class” University project of China Pharmaceutical University (CPU2018GF07, CPU2018PZQ19), Key research projects on modernization of traditional Chinese medicine (2018YFC1706900).

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Competing interests
The authors declare no competing interests.

Additional information
Supplementary information is available for this paper at https://doi.org/10.1038/s41598-020-60357-z.

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