

Research Article

The Value of Safflower Yellow Injection for the Treatment of Acute Cerebral Infarction: A Randomized Controlled Trial

Le-Jun Li,^{1,2} Yu-Mei Li,¹ Ben-Yu Qiao,¹ Shan Jiang,¹ Xin Li,¹
Hong-Ming Du,³ Peng-Cheng Han,² and Jiong Shi²

¹ Department of Neurology, Affiliated Lianyungang Hospital of Nanjing University of Chinese Medicine, No. 160 W. Chaoyang Road, Lianyungang 222000, China

² Barrow Neurological Institute, St. Joseph's Hospital and Medical Center, Phoenix, AZ 85013, USA

³ Changchun Sanzhen Industry Co., Ltd., Changchun 130000, China

Correspondence should be addressed to Le-Jun Li; lilej999@163.com

Received 8 August 2014; Revised 21 September 2014; Accepted 6 October 2014

Academic Editor: Feng-Qing Yang

Copyright © 2015 Le-Jun Li et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Background. Safflower Yellow Injection has been reported as a treatment for acute cerebral infarction in recent studies in China. However, there is a lack of availability of the evidence for the efficacy and safety of Safflower Yellow Injection for the treatment of acute ischemic stroke. So we investigated the effects of Safflower Yellow Injection for the treatment of acute cerebral infarction. **Method.** All subjects were randomly divided into Safflower Yellow Injection group given Safflower Yellow Injection (80 mg) and control group given placebo (0 mg) injection by intravenous drop once daily for 14 days. National Institute of Health Stroke Scale (NIHSS); hemorheological detection; coagulation function; and serum inflammatory markers, tumor necrosis factor- α (TNF- α), interleukin-1 β (IL-1 β), and interleukin-6 (IL-6), were used to investigate the effects before and 14 days after the treatment. **Results.** The scores of NIHSS were decreased on day 7 and day 14 after treatment. The hemorheological index of RBC deformation and RBC aggregation were significantly improved, prothrombin time (PT) increased, and fibrinogen (FIB) and TNF- α , IL-1 β , and IL-6 were decreased in patients treated with Safflower Yellow injection on day 14 after treatment ($P < 0.05$). **Conclusion.** Data suggests that Safflower Yellow Injection therapy may be beneficial for acute cerebral infarction.

1. Introduction

Stroke is a leading cause of death and disability worldwide [1]. Despite improvements in acute stroke care—stroke unit care, thrombolysis in appropriately selected patients and early and sustained antiplatelet therapy—many patients only make a partial or poor recovery after stroke and the major burden of stroke is chronic disability [2]. Therefore, there is a need for treatments that would further improve the efficacy. Clinical research in China, based on traditional Chinese medicine (TCM), has the potential of introducing new treatments for cerebral infarction.

Safflower Yellow Injection was certificated in China by the State Pharmaceutical Administration of China (SPAC) in 2002 (number 2002ZL0176) after being evaluated in clinical

trials. The constituent of Safflower Yellow Injection is Safflower Yellow, hydroxysafflor yellow A is a major component of Safflower Yellow [3]. The protective effect of hydroxysafflor yellow A in ischemic stroke has been investigated in previous studies [4], which have shown effects of antithrombosis [5, 6], anticoagulation [7], antioxidation [8], anti-inflammation [9], and anticalcium dysregulation [10].

Safflower Yellow Injection has been reported as a treatment for acute cerebral infarction in recent studies in China [11, 12]. However, there is a lack of good-quality and convincing supporting evidence for the efficacy and safety of Safflower Yellow Injection for the treatment of acute cerebral infarction [4]; no conclusions about efficacy or safety could be drawn [13]. Therefore, our study hypothesis was that patients who underwent a treatment with Safflower Yellow Injection

for acute cerebral infarction would have a better therapeutic effect compared to those who did not receive this therapy. So the aim of our study was to assess if giving Safflower Yellow Injection can improve effects of acute cerebral infarction patients in China.

2. Materials and Methods

2.1. Study Design. The study was a prospective, single-blinded, and randomized controlled trial. All eligible patients included in the study agree to participate and signed the informed consent form and the study procedures were approved by the Ethical Committee of the Affiliated Lianyungang Hospital of Nanjing University of Chinese Medicine. Full compliance with the Helsinki Declaration was applied during the study. Patients were recruited from the Neurology Department of the Affiliated Lianyungang Hospital of Nanjing University of Chinese Medicine between January 2004 and December 2012.

2.2. Inclusion and Exclusion Criteria. All patients meeting the World Health Organization's definition of ischemic stroke were enrolled during the study. Diagnostic criteria of acute cerebral infarction were diagnosed according to diagnostic criteria of various types of cerebrovascular disease [14] and China Guideline for Cerebrovascular Disease Prevention and Treatment. The main inclusion criteria of the trial were diagnosed ischemic stroke patients within 6 h to 14 days of onset; a score of 4–24 points on the National Institute of Health Stroke Scale (NIHSS); adults between 35 and 80 years old; and signed informed consent form. The main exclusion criteria were as follows: after 14 days from onset; a history of previous stroke; hemorrhagic stroke and transient ischemic attacks; other severe diseases such as heart or kidney failure, tumors, and gastrointestinal hemorrhage; and pregnant or lactating women.

2.3. Control and Intervention Groups. All patients of both groups were hospitalized for 14 days; we provide baseline treatment for all subjects to keep the respiratory tract unobstructed, preventing and treating aspiration pneumonia, monitoring and treating arrhythmia and ischemic heart disease, normalizing the blood pressure, and controlling blood sugar.

All patients were given conventional antiplatelet aggregation drugs (oral aspirin tablet, 100 mg, once daily) and neuroprotective agent (oral Piracetam tablet 400 mg, tid) therapy, based on conventional treatment; the Safflower Yellow Injection (80 mg) was given to the Safflower Yellow Injection group and placebo (0 mg) injection was given to the control group by intravenous drop once daily for 14 days.

The constituent of Safflower Yellow Injection is Safflower Yellow. The investigational drug Safflower Yellow Injection was supplied as Nankin freeze-dried powder. Each unit contains 80 mg Safflower Yellow. The placebo drug was supplied as blank colorless freeze-dried powder, where each unit contains no Safflower Yellow. Investigational drug and placebo were provided by Changchun Sanzhen Industry

Co., Ltd. The Safflower Yellow Injection group intravenously received 80 mg Safflower Yellow dissolved in 250 mL 0.9% sodium chloride or 5% glucose once per day. The control group was given a placebo. For two groups, the course of treatment lasted for 14 days.

2.4. Objectives and Outcome Measures

2.4.1. Measurement of NIHSS Score. The NIHSS was measured for stroke severity. The primary outcome measures were the differences in patients' scores on this scale among baseline, day 7, and day 14. The score was assessed by an independent clinical investigator.

2.4.2. Measurement of Hemorheological Detection. Blood samples were collected from ulnar vein before and after the treatment, the index of hemorheology included the whole blood viscosity, reduced viscosity, plasma viscosity, hematocrit, and the index of red cell deformity.

Keep 6 mL blood sample in an anticoagulation cuvette mixed with Heparin 100 μL , with LG-R-80 type blood viscosity tester and LG-B-190 type cytomorphosis/aggregation tester, at $37^\circ\text{C} \pm 0.1^\circ\text{C}$ and detect the whole blood hypsotomy coefficient of viscosity (shear rate 150 s^{-1}), hypotomy coefficient of viscosity (shear rate 5 s^{-1}), plasma viscosity, and index of red blood cell deformation and aggregation. The HCT determination used decigram method centrifugation in common temperature with rotary speed 3000 r/min for 30 min.

2.4.3. Measurement of Coagulation Function. Blood samples were collected from ulnar vein before and after the treatment, kept 6 mL blood sample in an anticoagulation cuvette mixed with Heparin 100 μL , centrifuged the sample immediately in common temperature with rotary speed 3000 r/min for 15 minutes and then collected serum and froze it at -80°C to measure. The values of prothrombin time (PT), activated partial thromboplastin time (APTT), fibrinogen (FIB), and international normalized ratio (INR) were detected by using blood coagulation method before and after the treatment.

2.4.4. Measurement of Serum Inflammatory Markers $\text{TNF-}\alpha$, $\text{IL-1}\beta$, and IL-6 . Peripheral venous blood (10 mL) was collected into sampling tubes without EDTA. Blood was centrifuged at 3500 rpm for 5 minutes to separate the serum. Serum was stored at -80°C until analysis. All assays were performed in a blinded fashion on coded samples. Serum levels of $\text{TNF-}\alpha$, $\text{IL-1}\beta$, and IL-6 were measured in duplicate by a sandwich-type enzyme-linked immunosorbent assay (ELISA) technique by using kits from R&D Systems, $\text{TNF-}\alpha$ DTA00C (range of detection: 15.6–1,000 pg/mL), $\text{IL-1}\beta$ DLB50 (range of detection: 3.9–250 pg/mL), and IL-6 D6050.

2.5. Statistical Analysis. Statistical analysis was performed using SPSS 12.0 statistics software. Data was presented as mean \pm S.D.; baseline variables were analyzed using student's *t*-test. For efficacy variables, they were analyzed using analysis of covariance (ANOVA); comparisons were made between

the two groups at baseline, day 7, and day 14. The two-group *t*-test was used separately for each comparison. A level of $P < 0.05$ was considered to be statistically significant.

3. Results

3.1. Baseline Characteristics. 150 patients were approached and screened for their eligibility to enroll in this study. 108 patients consented and were eligible for the study and signed their informed consent for participation. Patients were randomized by a central stochastic system (2:1) to either Safflower Yellow Injection group ($n = 72$) or control group ($n = 36$). Seven patients were rejected during the treatment period for not meeting the inclusion criteria (four from the Safflower Yellow Injection group and three from the control group) and another four patients dropped out for adverse events (two from the Safflower Yellow Injection group and two from the control group). Ninety-seven patients completed the trial, 66 patients in the Safflower Yellow Injection group and 31 patients in the control group (see Figure 1). The characteristics of the Safflower Yellow Injection and control groups are not significantly different at baseline and are given in Table 1.

3.2. Efficacy Results. The scores of NIHSS are all differently reduced after treatment for 7 and 14 days. This improvement becomes more apparent with time. After subtracting the baseline effect, the between-group total score is statistically significant ($P < 0.01$) after treatment for 7 and 14 d.

There were no significant differences in the values of hemorheological index between the two groups before the treatment ($P > 0.05$). Safflower Yellow Injection group hemorheological index of RBC deformation and index of RBC aggregation; there were significant differences before and after the treatment ($P < 0.05$). In the control group, only the whole blood viscosity was significantly different before and after the treatment ($P < 0.05$); the remaining hemorheological indices were not significantly different before and after treatment. Safflower Yellow Injection group's whole blood viscosity and plasma viscosity were significantly different after treatment between the two groups ($P < 0.05$). Results are listed in Table 2. The reason may be that Safflower Yellow Injection can improve the values of hemorheological index.

There were no significant differences in the PT, APTT, INR, and FIB values between the two groups before the treatment ($P > 0.05$). The value of PT increased in the Safflower Yellow Injection group after treatment ($P < 0.05$); there was significant difference between the two groups ($P < 0.05$). After the treatment, the values of FIB decreased in the Safflower Yellow Injection group after treatment; there was significant difference between the two groups ($P < 0.05$). Results are listed in Table 3. The reason may be that Safflower Yellow Injection can inhibit both intrinsic and extrinsic coagulations.

There were no significant differences in the serum levels of TNF- α , IL-1 β , and IL-6 between the two groups before the treatment ($P > 0.05$). The serum levels of TNF- α , IL-1 β , and IL-6 in the Safflower Yellow Injection group decreased

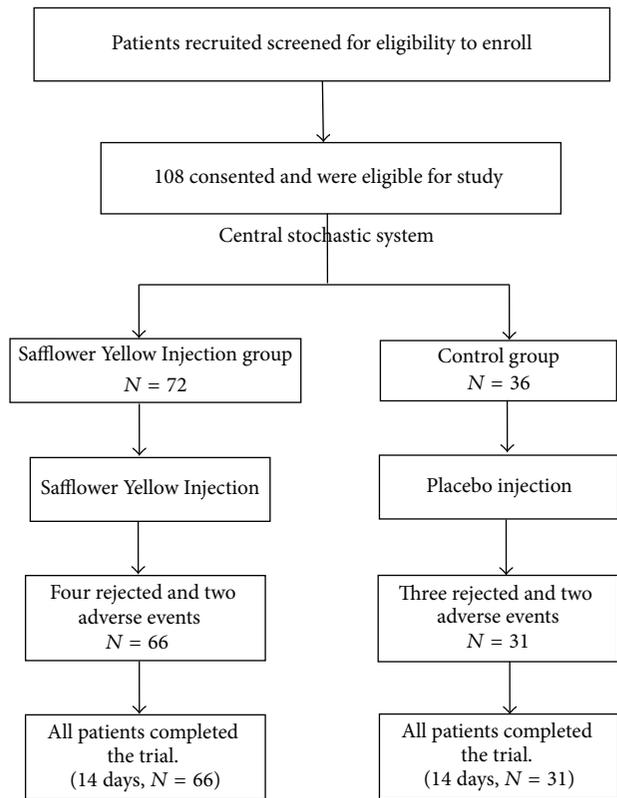


FIGURE 1: Study design and CONSORT diagram showing the flow of participants.

after treatment ($P < 0.05$); there were significant differences between the two groups ($P < 0.05$). Results are listed in Table 4. The reason may be that Safflower Yellow Injection has anti-inflammatory effects.

3.3. Safety and Tolerability. Safflower Yellow Injection for the treatment acute cerebral infarction was well tolerated. Four adverse events were reported during the whole trail (two in the Safflower Yellow Injection group and two from the control group). In the Safflower Yellow Injection group, one patient suffered a transfusion reaction and the other was diagnosed of poststroke depression. Two patients in the control group suffered a recurrence of stroke. All adverse events were considered not to be related to the Safflower Yellow Injection.

4. Discussion

Stroke is a sudden onset and acute cerebral vascular disease commonly characterized by focal neurological deficit. It is the most common neurological disease with high incidence, morbidity, mortality, and recurrence. The disease is more common in the elderly, together with myocardial infarction and cancer; these are the three major causes for senile fatality [15].

This clinical study showed that Safflower Yellow Injection is safe and effective for acute cerebral infarction. It effectively

TABLE 1: Baseline characteristics of study patients.

Characteristics <i>n</i> = 97	Control group <i>n</i> = 31	Safflower Yellow Injection group <i>n</i> = 66	<i>P</i> value
Male	9	28	0.595
Mean age and range, years	58.84 ± 10.18 (30–70)	59.47 ± 10.92 (31–70)	0.564
Body height/cm	Male	170.00 ± 6.69	0.573
	Female	159.19 ± 4.82	0.554
Body weight/kg	Male	67.56 ± 9.48	0.563
	Female	58.73 ± 8.18	0.612
Weight index/(kg/m ²)	23.19 ± 2.49	23.79 ± 2.76	0.963
Disease time/d	5.00 ± 3.55	3.91 ± 2.77	0.925
NIH stroke scale score	7.65 ± 2.32	8.53 ± 3.30	0.556

Values are mean ± S.D.; no significant changes between control group and Safflower Yellow Injection group.

TABLE 2: Comparison of hemorheological index before and after treatment between 2 groups ($\bar{x} \pm s$).

Variables	Control group <i>n</i> = 31		Safflower Yellow Injection group <i>n</i> = 66	
	Pretreatment	Posttreatment	Pretreatment	Posttreatment
Whole blood viscosity (mPa·s/150 s ⁻¹)	5.95 ± 0.80	5.34 ± 0.83 [#]	5.92 ± 0.82	5.14 ± 0.85 ^{##}
Whole blood viscosity (mPa·s/5 s ⁻¹)	12.66 ± 2.52	12.57 ± 2.24	12.69 ± 2.64	10.05 ± 2.48 ^{##}
Plasma viscosity (mPa·s)	1.82 ± 0.23	1.88 ± 0.26	1.78 ± 0.21	1.42 ± 0.23 ^{##}
HCT (%)	49.76 ± 4.80	47.64 ± 4.95	48.53 ± 5.30	47.61 ± 4.77
Index of RBC deformation	0.51 ± 0.01	0.50 ± 0.01	0.51 ± 0.07	0.59 ± 0.03 [*]
Index of RBC aggregation	1.21 ± 0.16	1.36 ± 0.19	1.38 ± 0.19	1.21 ± 0.16 [*]

Data are expressed as mean ± S.D., **P* < 0.05, compared with the pretreatment, [#]*P* < 0.05, compared with control group.

TABLE 3: Changes of coagulation and fibrolysis indexes in the patients before and after treatment between 2 groups ($\bar{x} \pm s$).

Variables	Control group <i>n</i> = 31		Safflower Yellow Injection group <i>n</i> = 66	
	Pretreatment	Posttreatment	Pretreatment	Posttreatment
PT (s)	10.64 ± 0.58	10.85 ± 0.57	11.02 ± 0.65 [#]	13.38 ± 0.71 ^{##}
APTT (s)	29.43 ± 4.76	29.21 ± 4.83	29.84 ± 4.81	29.06 ± 5.02
INR	1.04 ± 0.13	0.94 ± 0.12	1.04 ± 0.15	0.94 ± 0.16
FIB (g/L)	3.35 ± 0.61	3.54 ± 0.70	3.57 ± 0.73	2.07 ± 0.19 ^{##}

Data are expressed as mean ± S.D., **P* < 0.05, compared with the pretreatment, [#]*P* < 0.05, compared with control group.

TABLE 4: Comparison of serum levels of TNF- α , IL-1 β , and IL-6 before and after treatment between 2 groups ($\bar{x} \pm s$).

Variables	Control group <i>n</i> = 31		Safflower Yellow Injection group <i>n</i> = 66	
	Pretreatment	Posttreatment	Pretreatment	Posttreatment
TNF- α	39.45 ± 5.62	40.24 ± 1.55	39.82 ± 4.71	30.85 ± 3.10 ^{##}
IL-1 β	44.16 ± 4.32	40.07 ± 1.44	44.69 ± 4.24	28.85 ± 1.68 ^{##}
IL-6	669.92 ± 61.93	711.28 ± 31.86	689.88 ± 48.92	558.82 ± 38.03 ^{##}

Data are expressed as mean ± S.D., **P* < 0.05, compared with the pretreatment, [#]*P* < 0.05, compared with control group.

improved the stroke symptoms and significantly improved the patient's life competency. The NIHSS is widely used in nearly all large clinical stroke trials to document baseline and outcome severity of neurological impairment [16]. In our study, we found that the scores of NIHSS were significantly decreased in patients treated with Safflower Yellow on day 7

and day 14 after treatment. These results suggest that Safflower Yellow is beneficial in reducing the severity of stroke.

Hemorheological detection is important in the course of disease, including nosogenesis, development, prognosis, and turnover as well as curative effect assessment, especially for prevention of cardiocerebrovascular disease. Brain

microcirculation slowdown or stagnation is the main pathological change at the beginning of cerebral ischemia [17]. In our study, we found that the hemorheological index of RBC deformation and RBC aggregation were significantly improved in patients treated with Safflower Yellow Injection on day 14 after treatment. These results suggest that Safflower Yellow Injection has protective effect; the reason may be that Safflower Yellow Injection can reduce whole blood viscosity and plasma viscosity, improve index of RBC deformation, reduce index of RBC aggregation, improve microcirculation, and improve brain following ischemic reperfusion in the acute cerebral infarction; the protective effect of hydroxysafflower yellow A in ischemic stroke has been investigated in previous studies [18].

Coagulation function detection is important in the course of disease, including nosogenesis and curative effect assessment of acute cerebral infarction. In our study, we found that the value of PT increased in the Safflower Yellow Injection group after treatment and the values of FIB decreased in the Safflower Yellow Injection group after treatment, and there were significant differences between the two groups. These results suggest that Safflower Yellow Injection can inhibit both intrinsic and extrinsic coagulations, significantly the prothrombin time and clotting time, and reduce platelet adhesion, thrombosis, and fibrin cross-linking process. Recent studies show that Safflower Yellow inhibits platelet activation factor-induced platelet aggregation, antagonizes platelet activation factor-induced capillary permeability [19, 20], and protects against antithrombin III injuries originated from free radicals [21].

Researchers have demonstrated that the following brain insult cytokine levels are elevated as a result of increased production from inflammatory cells [22] with IL-1, IL-6, and tumor necrosis factor- α (TNF- α) and being the most studied for stroke [23, 24]. IL-1 β and TNF- α have been associated with exacerbation of injury in stroke while IL-6 has been found to be neuroprotective [25]. In our study, we found that the serum levels of TNF- α , IL-1 β , and IL-6 were significantly decreased in patients treated with Safflower Yellow Injection on day 14 after treatment. These results suggest that Safflower Yellow Injection is beneficial in reducing the serum levels of TNF- α , IL-1 β , and IL-6 after treatment. The reason may be that Safflower Yellow Injection has anti-inflammatory effects; it is considered to be the main source of antioxidant defense in the brain following ischemic reperfusion; the protective effect of hydroxysafflower yellow A in ischemic stroke has been investigated in previous studies [12].

Our data suggest that Safflower Yellow Injection therapy may be beneficial for acute cerebral infarction. In the Chinese studies, Safflower Yellow Injection exhibits a favorable safety profile; there were no serious adverse events recorded and only 2 cases adverse events. This low rate of adverse events may be due to a combination of the fact that the patients were recruited during their recovery phase when their clinical condition had stabilized and due to the method of collection of adverse events in China.

However, there are still some limitations in our study. For example, the size (108 subjects) was not sufficient. The period

of observation lasted for only 14 days. To further studies with larger sample size and long-term observations are required. Therefore, we could not rule out the expectation bias because neither the patients nor the therapists were blinded. Some future double-blinded studies would be feasible to be carried out in the future by using Safflower Yellow Injection.

5. Conclusions

In summary, our data suggest that Safflower Yellow Injection therapy may be beneficial for acute cerebral infarction. The results presented here invite for further studies with larger sample size, long-term observation, and strict blinding to confirm the efficacy of Safflower Yellow Injection.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

Acknowledgments

The authors would like to thank the staff of Department of Neurology of the Affiliated Lianyungang Hospital of Nanjing University of Chinese Medicine for their assistance in recruiting patients for the study. This study was financed by Jiangsu Health International Exchange Program, China.

References

- [1] C. D. Mathers and D. Loncar, "Projections of global mortality and burden of disease from 2002 to 2030," *PLoS Medicine*, vol. 3, no. 11, article e442, 2006.
- [2] C. D. A. Wolfe, "The impact of stroke," *British Medical Bulletin*, vol. 56, no. 2, pp. 275–286, 2000.
- [3] Z. F. Yang, Q. B. Mei, and Y. P. Jiang, "The effective components and pharmacological effects of safflower," *Xi Bei Yao Xue Za Zhi*, vol. 16, no. 2, pp. 131–133, 2001 (Chinese).
- [4] S. Fan, N. Lin, G. Shan, P. Zuo, and L. Cui, "Safflower yellow for acute ischemic stroke: a systematic review of randomized controlled trials," *Complementary Therapies in Medicine*, vol. 22, no. 2, pp. 354–361, 2014.
- [5] B.-X. Zang, M. Jin, N. Si, Y. Zhang, W. Wu, and Y.-Z. Piao, "Antagonistic effect of hydroxysafflower yellow a on the platelet activating factor receptor," *Yaoxue Xuebao*, vol. 37, no. 9, pp. 696–699, 2002 (Chinese).
- [6] H. B. Zhu, L. Zhang, Z. H. Wang et al., "Therapeutic effects of hydroxysafflower yellow A on focal cerebral ischemic injury in rats and its primary mechanisms," *Journal of Asian Natural Products Research*, vol. 7, no. 4, pp. 607–613, 2005.
- [7] X. Sun, X. Wei, S. Qu, Y. Zhao, and X. Zhang, "Hydroxysafflower yellow A suppresses thrombin generation and inflammatory responses following focal cerebral ischemia-reperfusion in rats," *Bioorganic & Medicinal Chemistry Letters*, vol. 20, no. 14, pp. 4120–4124, 2010.
- [8] L. Sun, L. Yang, Y.-W. Xu et al., "Neuroprotection of hydroxysafflower yellow A in the transient focal ischemia: inhibition of protein oxidation/nitration, 12/15-lipoxygenase and bloodbrain barrier disruption," *Brain Research*, vol. 1473, no. 9, pp. 227–235, 2012.

- [9] S.-Y. Ye and W.-Y. Gao, "Hydroxysafflor yellow A protects neuron against hypoxia injury and suppresses inflammatory responses following focal ischemia reperfusion in rats," *Archives of Pharmacological Research*, vol. 31, no. 8, pp. 1010–1015, 2008.
- [10] J.-W. Tian, F.-H. Fu, W.-L. Jiang, C.-Y. Wang, F. Sun, and T.-P. Zhang, "Protective effect of hydroxysafflor yellow A against rat cortex mitochondrial injuries induced by cerebral ischemia," *Yaoxue Xuebao*, vol. 39, no. 10, pp. 774–777, 2004 (Chinese).
- [11] Z. L. Ma, B. X. Wu, D. M. Cheng et al., "Clinic observation of the curative effect of safflower yellow in the treatment of acute cerebral infarction," *Chinese Journal of Practical Nervous Disease*, vol. 14, no. 24, pp. 30–31, 2011 (Chinese).
- [12] B. Ye, W. J. Li, L. Zheng, and R. Liu, "Effect of safflower yellow pigment on blood rheology and cerebral hemodynamics in patients with acute cerebral infarction," *Chinese Journal of General Practice*, vol. 10, no. 2, pp. 219–221, 2012 (Chinese).
- [13] B. Wu, M. Liu, H. Liu et al., "Meta-analysis of traditional Chinese patent medicine for ischemic stroke," *Stroke*, vol. 38, no. 6, pp. 1973–1979, 2007.
- [14] Chinese Neuroscience Society and Chinese Neurosurgical Society, "Diagnostic criteria of various types of cerebrovascular disease," *Zhong Hua Shen Jing Ke Za Zhi*, vol. 29, no. 6, pp. 89–90, 1996 (Chinese).
- [15] J. Wu, J. P. Jia, and L. Y. Cui, *Neurology*, People's Health Publishing Company, Beijing, China, 2005, (Chinese).
- [16] L. B. Goldstein and G. P. Samsa, "Reliability of the National Institutes of Health Stroke Scale. Extension to non-neurologists in the context of a clinical trial," *Stroke*, vol. 28, no. 2, pp. 307–310, 1997.
- [17] A. Bruno, J. P. McConnell, S. N. Cohen et al., "Serial urinary 11-dehydrothromboxane B₂, aspirin dose, and vascular events in blacks after recent cerebral infarction," *Stroke*, vol. 35, no. 3, pp. 727–730, 2004.
- [18] I. Velcheva, E. Titianova, and N. Antonova, "Evaluation of the hemorheological and neurosonographic relationship in patients with cerebrovascular diseases," *Clinical Hemorheology and Microcirculation*, vol. 30, no. 3-4, pp. 373–380, 2004.
- [19] W. Wu, M. Jin, and S. D. Yang, "Safflower yellow inhibition of platelet-activating factor due to the proliferation of capillary permeability," *Cardiopulmonary Disease Journal*, vol. 18, no. 4, pp. 281–283, 1999.
- [20] W. M. Kuebler, Y. Yang, R. Samapati, and S. Uhlig, "Vascular barrier regulation by PAF, ceramide, caveolae, and NO—an intricate signaling network with discrepant effects in the pulmonary and systemic vasculature," *Cellular Physiology and Biochemistry*, vol. 26, no. 1, pp. 29–40, 2010.
- [21] J. W. Li, "Safflower yellow plasminogen activator inhibiting agents dose levels in rabbit," *Chinese Herbal Medicine*, vol. 30, no. 2, pp. 128–129, 1999 (Chinese).
- [22] T. Sairanen, O. Carpén, M.-L. Karjalainen-Lindsberg et al., "Evolution of cerebral tumor necrosis factor- α production during human ischemic stroke," *Stroke*, vol. 32, no. 8, pp. 1750–1757, 2001.
- [23] H. S. Han and M. A. Yenari, "Cellular targets of brain inflammation in stroke," *Current Opinion in Investigational Drugs*, vol. 4, no. 5, pp. 522–529, 2003.
- [24] A. Durukan and T. Tatlisumak, "Acute ischemic stroke: overview of major experimental rodent models, pathophysiology, and therapy of focal cerebral ischemia," *Pharmacology Biochemistry and Behavior*, vol. 87, no. 1, pp. 179–197, 2007.
- [25] S. M. Allan and N. J. Rothwell, "Cytokines and acute neurodegeneration," *Nature Reviews Neuroscience*, vol. 2, no. 10, pp. 734–744, 2001.



Hindawi
Submit your manuscripts at
<http://www.hindawi.com>

