

Safflower yellow pigment and Sanqi *Panax notoginseng* in the treatment of acute cerebral infarction: a systematic review, meta-analysis, and cost-effectiveness analysis

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Background: Sanqi *Panax notoginseng* injection and safflower yellow injection were Chinese traditional medicine injections for the treatment of cardiovascular diseases and were used to treat acute cerebral infarction patients in public hospital widely. The aim of this study was to compare and analyze the published reports of efficacy and safety of Sanqi *Panax notoginseng* injection and safflower yellow injection for the treatment of acute cerebral infarction. The cost-effectiveness of these drug formulations was also evaluated.

Methods: China National Knowledge Infrastructure (CNKI), Wanfang, SinoMed, VIP, PubMed, Embase, and the Chinese Biomedical Literature (CBM) were searched with the restrictions keywords in Chinese and English between 2006 and 2019 to obtain RCTs. A meta-analysis and a meta-regression analysis were undertaken in Reviewer Manager 5.3 software to compare the efficacy and safety of Sanqi *Panax notoginseng* and safflower yellow injection. This study used a decision tree model to analyze the cost-effectiveness of the two treatments. The TreeAge Pro software was used to comprehensively evaluate the economics of these medications.

Results: Twelve papers were all randomized controlled trials (RCTs) in which Sanqi *Panax notoginseng* injection was applied in the control group, while safflower yellow injection was applied in the experimental group and the quality of them were good. The results of the 12 papers were compared, and the total effective rate of the treatment group (91.18%) was significant and showed no significant difference with the control group (74.83%) (RR =1.24, 95% CI: 1.19, 1.30, P<0.00001). From the perspective of pharmacoeconomics, compared with Sanqi *Panax notoginseng* group, the ICER of safflower yellow injection is 3,885.75 RMB. The sensitivity analysis results were consistent with the basic analysis results, indicating that the basic analysis results were relatively stable.

Conclusions: Comparing with Sanqi *Panax notoginseng* injection, safflower yellow injection and related combination therapy can improve the total effective rate and are safer with fewer adverse reactions. It is also more cost-effective than the use of Sanqi *Panax notoginseng* injection.

Keywords: Safflower yellow pigment; safflower injection; cerebral infarction; meta-analysis; cost-effectiveness analysis

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Introduction

Cerebral infarction, also known as ischemic stroke, is a common disease in clinical practice. It is caused by local lesions in the cerebral vessel wall, leading to vasospasm, stenosis, or occlusion, thus causing impaired blood supply to the brain, irreversible damage to the local brain tissue, and ultimately ischemic and hypoxic brain tissue necrosis (1-3). According to the statistics of the World Health Organization (WHO), in 2010 and 2016, cerebral infarction was the second leading cause of death globally, next only to ischemic heart disease. In China, cerebral infarction has been the first leading cause of death, with more than 1.5 million new cases each year (4). The Institute of Health and Economics of the Chinese Ministry of Health has reported that the social and economic burden caused by cerebrovascular disease in China is as much as 20 billion yuan per year, and the incidence of cerebral infarction is the most prevalent subtype, so the disease burden it brings is extreme (5). Because most cerebral infarction have a sudden onset and result in a high disability rate, most patients lack sufficient psychological preparation and develop negative emotions such as fear and anxiety, contributing to a psychological burden (6). In sum, cerebral infarction put extra pressure and burden on patients, their families, and the whole society.

Previous studies have shown that Sangi Panax notoginseng injection, with notoginoside as the main component, has positive therapeutic effects on cerebral infarction (7). It can also significantly improve the neurological function and quality of life of patients. Safflower yellow injection is mainly composed of the yellow pigment extracted from safflower (Carthamus tinctorius). Previous studies have shown that safflower yellow injection significantly dilates blood vessels, improves microcirculation and anticoagulation and reduces blood viscosity, with a beneficial effect for the treatment of coronary heart disease, myocardial ischemia, hyperviscosity, and ischemic cerebrovascular disease (8-14). At present, Sanqi Panax notoginseng and safflower yellow pigment have been all clinically applied to the treatment of cerebral infarction in public hospitals because they were included in China Medical Insurance Directory. However, there is little information reported on the pharmacoeconomic evaluation

and the comparison of efficacy and safety between them. The aim of this study was to review the current evidence on the therapeutic effects of safflower yellow injection and Sanqi *Panax notoginseng* on cerebral infarction. A cost-effectiveness analysis was constructed based on the decision tree model to evaluate the economics of the two drug regimens. The best clinical treatment plan in terms of effectiveness, safety, and economy must be determined to provide guidance for clinical rational drug use, rational resource allocation and medical cost savings. We present the following article in accordance with the PRISMA reporting checklist (available at https://dx.doi.org/10.21037/atm-21-782).

Methods

The PRISMA guidelines (15) were used for designing and reporting this study, and this study was in accordance with the ethical guidelines of the Declaration of Helsinki (as revised in 2013) (16). Our PROSPERO registration number is CRD42020167326.

Search strategy

We searched Chinese database such as China National Knowledge Infrastructure (CNKI), Wanfang, SinoMed, and the Chinese Biomedical Literature (CBM); English database such as PubMed, Embase and web of science from January 2006 to January 2019. The keywords included in the search were "safflower yellow", "acute cerebral infarction", and "Sanqi *Panax notoginseng*". The references cited by the included studies were also evaluated to locate more relevant studies. The ClinicalTrial.gov database was also polled to include human trial results that had not been reported elsewhere. Grey literatures and ongoing trials were searched. We only searched studies in Chinese and English.

Study selection

The participants of these studies were patients with acute cerebral infarction. Safflower yellow pigment including safflower yellow injection and safflower injection were the intervention of experiment group. Sanqi *Panax*

notoginseng injection was the comparison intervention. The route of administration was intravenous drip. The outcome indicator was the total effective rate and National Institutes of Health Stroke Scale (NIHSS) score. The total effective rate is defined based on the NIHSS score: (I) significantly effective: the clinical symptoms of the patient completely disappeared. Loss of function score is greater than 90%, and no disability occurs: (II) effective: the patient's clinical symptoms have been significantly improved, and the vital signs also have more significant changes, the loss of function score is less than 50%: the degree of disability is less than 3; (III) invalid: the patient's clinical symptoms and vital signs have not changed, and the condition even worsened.

Study types were limited to RCTs and retrospective trials. Animal experiments and case studies were excluded. Studies with incomplete data and duplicate data were excluded.

Data extraction and quality assessment

Two authors independently assessed the quality of studies for bias analysis using the Cochrane Handbook for Systematic Reviewers version 5.1.0 risk of bias assessment tool for RCTs (17). The following items were assessed: (I) whether the random method is correct; (II) whether allocation concealment is achieved; (III) whether blinding is used (implementation bias); (IV) blinding of outcome evaluators (measurement bias); (V) completeness of outcome data (follow-up bias); (VI) selective reporting (reporting bias); and (VII) other sources of bias. Each aspect of the content had three levels, including "high", "unclear", and "low". In the cases of disagreement, a third party adjudicated.

Statistical analysis

Statistical analysis was undertaken in Review Manager 5.3 (Cochrane Collaboration, Copenhagen, Denmark). Odds ratio (OR) was used as the effect indicator for enumeration data, and mean ($\bar{x} \pm s$) was used as the effect indicator for measurement data. The point estimate and 95% confidence interval (CI) were given for each effect size. Heterogeneity among the results of the included studies was tested using the χ^2 test, with P<0.05 as the test level, and the magnitude of heterogeneity was quantitatively determined in combination with I². If there was no significant heterogeneity among studies (P>0.05, I^2 <50%), the fixed effects model was selected for effect size combination; if there was heterogeneity among studies (P<0.05, I²>50%), the random effects model was used for effect size combination. Publication bias was assessed using funnel plots. At the same time, the Egger and Begger values were calculated using STATA14.0 to judge whether there was bias. P>0.05 was taken as the test level. If P>0.05, it was considered that the meta-analysis was unbiased, otherwise there was bias.

In the evaluation of pharmacoeconomics, costeffectiveness analysis was used to calculate the incremental cost-effectiveness ratio; single-factor sensitivity analysis was used to draw the cyclone chart. Monte Carlo simulation was used for probability sensitivity analysis, and then a cost-effectiveness curve was drawn (18). In this paper, TreeAge2011 software was used to build a decision tree model for cost-effectiveness analysis and sensitivity analysis.

Pharmacoeconomic evaluation

Research perspective

This study used cost-effectiveness analysis for pharmacoeconomic comparison, considering the cost from the perspective of patients and calculating the treatment efficiency for patients with cerebral infarction. The years of improvement in quality of life, indirect costs of treatment, and hidden costs that cause inconvenience to patients were not considered, because there is no standard available for reference. Therefore, the direct cost of the drug itself, that is, the price per use, was considered to be the cost. In addition, the effective components of safflower yellow injection and safflower injection in this study are the same, although they are produced by different manufacturers. These data were kept separate in the above meta-analysis, and were been evaluated differently in the cost calculation. The decision indicator of the cost-effectiveness analysis was the incremental cost-effectiveness ratio, and the smaller ratio is indicative that the pharmacoeconomic advantage is better.

Decision tree model

Model structure: this study used a decision tree model to analyze the cost-effectiveness of Safflower yellow injection and Sanqi *Panax notoginseng* injection included in the meta-analysis for the treatment of cerebral infarction diseases. The meta-analysis method was used to obtain the effectiveness and safety indicators, and comprehensively

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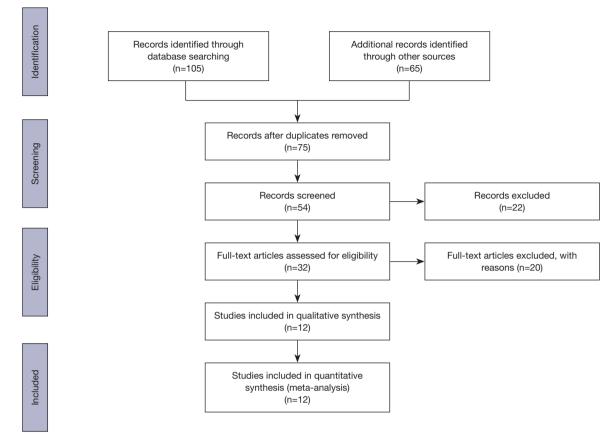


Figure 1 Flow diagram of the literature review.

evaluate the economics of safflower yellow and Sanqi *Panax notoginseng*. The model mainly evaluated the short-term economics of safflower yellow and Sanqi *Panax notoginseng* injection. The time span of the model simulation was a course of treatment (14 days).

Effectiveness: in this study, the effective rate of treatment for patients with cerebral infarction was the efficiency index. According to the weight of each study obtained from meta-analysis, the treatment efficiency of cerebral infarction patients in each study was weighted, and the effectiveness included in the pharmacoeconomic evaluation was obtained.

Effectiveness index: judging whether it is effective according to the total effective rate, effective was counted as 1, invalid or ineffective was counted as 0.

Cost: this study analyzed cost from the perspective of patients. Since the goal of the patient is to obtain the best prevention and diagnosis results with the least personal expenditure and minimal costs, only the cost to the patient and their family was calculated (19).

Results

Literature search results

Literature search and screening results

Initially, 170 literature reports were identified. After excluding duplicate literatures, 75 literatures were identified. Unrelated papers such as, animal experiments, and case reports were excluded. A refined set of 32 clinical studies on safflower injection for cerebral infarction were remained. After further reading of the full text of each study, all literatures with incomplete data and duplicate data as well as unrelated outcome indicators were excluded. Finally, 12 Chinese literatures reports were included in the study. The literature screening process is shown in *Figure 1*.

Literature search results and quality evaluation

The 12 literature reports of RCTs that were included had a total of 1,425 patients, among whom 714 were in the

experimental group and 711 were in the control group. The literature was evaluated using the Cochrane System Evaluator's Manual 5.1.0 RCT bias risk assessment tool. After evaluating, all studies achieved a score no less than 3, which indicating that all studies had good methodological quality. The summarized content for each study (13,20-29) is shown in *Table 1*.

Meta-analysis results

Total effective rate of safflower yellow pigment

The 12 studies (13,20-29) that reported the effective rate were analyzed under a random effect model. Meta-analysis showed a significantly higher overall effective rate in safflower yellow group compared with that in the control group (RR =1.24, 95% CI: 1.19, 1.30, P<0.00001) (*Figure 2*) (30). There was no evidence of heterogeneity between these studies (P=0.009, I^2 =56%).

Meta-regression analysis

According to the results of the previous meta-analysis, heterogeneity exists among some studies (13,20-29) of safflower yellow in the treatment of cerebral infarction. In order to search for the causes of heterogeneity and the factors affecting the treatment effect, such as duration of treatment, age, gender, number of trials, and whether to use the blind method, a multivariate meta-regression analysis was performed for the age and the type of injection. The results are shown in *Table 2*.

It can be seen that the between-study heterogeneity tau^2 =0.001778 is 0.006322 less than the previous 0.0081 after adding the two variables of age and type of injection. This indicates that different age and different types of injection production can explain 78.05% of the heterogeneity. Therefore, we considered the subgroup analysis of safflower yellow injection and safflower injection, which could effectively explain the heterogeneity and obtain reasonable conclusions.

Total effective rate of safflower yellow injection

Seven studies that reported the effective rate (21,22,24-28) were analyzed under a fixed effect model. Meta-analysis showed a significantly higher effective rate in safflower yellow group compared with that in control group (OR =4.33, 95% CI: 2.67, 7.05, P<0.00001) (*Figure 3*). There was no evidence of heterogeneity between these studies (P=0.68, I²=0).

Total effective rate of safflower injection

Five studies that reported the effective rate (13,20,23,29,30) were analyzed under a random effect model. Meta-analysis showed a significantly higher overall effective rate in safflower injection group compared with that in control group (OR =6.00, 95% CI: 3.71, 9.72, P<0.00001) (*Figure 4*). There was no evidence of heterogeneity between these studies (P=0.03, I^2 =62%).

Detection bias

The funnel plot of the total effective rate evaluated in the included papers (13,20-29) was analyzed (*Figure 5*). The funnel plot was essentially symmetric, suggesting absence of publication bias (Egger P=0.223; Begger P=0.2441).

Adverse reactions

Six studies reported that no adverse reactions were observed (20-25); one did not mention adverse reactions (30); and five reported the occurrence of adverse reactions (26-30). The details of the inclusion of adverse reactions can be seen in *Table 3*. The above results suggested that safflower yellow injection and Sanqi *Panax notoginseng* injection had none or mostly mild adverse effects on patients.

Pharmacoeconomic analysis

Cost calculation

The cost variable for this paper was calculated from the perspective of patients. Drug prices understandably fluctuate throughout time, so the retail price of the therapy was chosen closest to the time of the study. Cost-effectiveness analysis does not distinguish anything based on the severity of disease. Direct medical cost, direct non-medical cost and indirect cost are the average cost of each disease state (mild, moderate and severe) (31). In this study, it was assumed that the direct nonmedical cost and indirect cost of the patients in each group were the same, respectively, and the difference in the total cost was caused by the direct medical cost. In addition, due to the similar costs of hospitalization and drug administration in direct medical costs, this study only considered the cost of the drug itself. When the treatment time of the disease exceeds one year, the cost should be discounted. According to the recommendation of the international pharmacoeconomic study, a 5% discount rate was adopted for cost data and uniformly discounts were applied to 2020.

In this study, the investigation of drug prices was

Research ID	Number of cases (E/C)	Experimental group medication	Control group medication	Follow up date	Efficacy index	Cochrane remark	Document type
Gao JH, 2015	46/46	Safflower injection 100 mg + Sodium Ozagrel 80 mg	Sanqi <i>Panax notoginseng</i> 400 mg + Sodium Ozagrel 80 mg	14 d	Neurological deficit score	m	RCT
Peng W, 2017	23/23	Safflower injection 20 mL + Sanqi <i>Panax notoginseng</i> 400 mg	Sanqi <i>Panax notoginseng</i> 400 mg	14 d	Neurological deficit score	S	RCT
Kong LY, 2006	47/48	Safflower injection 20 mL	Sanqi <i>Panax notoginseng</i> 0.5 g	14 d	Neurological deficit score	ო	RCT
Gao JH, 2014	52/52	Safflower yellow pigment 150 mg + Yuriklin 0.15 PNA	Sanqi <i>Panax notoginseng</i> 400 mg + Yuriklin 0.15 PNA	14 d	Neurological deficit score	ю	RCT
Cheng QH, 2017	80/70	Safflower yellow pigment 100 mg + Sodium Ozagrel 80 mg	Sanqi <i>Panax notoginseng</i> 40 mg	15 d	Neurological deficit score	£	Retrospective analysis
Lv QX, 2015	15/15	Safflower yellow pigment 100 mg + Sodium Ozagrel 80 mg + conventional treatment	Sanqi <i>Panax notoginseng</i> 40 mg + conventional treatment	14 d	Neurological deficit score	Ŋ	RCT
Wen ZP, 2014	32/32	Safflower yellow pigment 100 mL + Sodium Ozagrel 80 mg	Sanqi <i>Panax notoginseng</i> 40 mg	14 d	Neurological deficit score	ი	RCT
Xing ZG, 2018	02/02	Safflower Injection 20 mL + Sodium Ozagrel 160 mg	Xueshuantong 500 mg + low molecular dextrose 500 mL	14 d	Neurological deficit score	ი	RCT
Miao MJ, 2014	30/30	Safflower yellow pigment 100 mL	Sanqi <i>Panax notoginseng</i> 400 mg	14 d	Neurological deficit score	£	RCT
Wang WL, 2012	120/120	Safflower Injection 20 mL + Sodium Ozagrel 160 mg	Sanqi <i>Panax notoginseng</i> 400 mg + Iow molecular dextrose 500 mL	14 d	Neurological deficit score	ი	RCT
Jin YQ, 2013	140/140	Safflower Injection 20 mL + Sodium Ozagrel 160 mg	Sanqi <i>Panax notoginseng</i> 400 mg + Iow molecular dextrose 500 mL	14 d	Neurological deficit score	4	RCT
Liu J, 2016	59/65	Safflower yellow pigment 25 mL	Sanqi <i>Panax notoginseng</i> 400 mg	14 d	Neurological deficit score	с	Retrospective analysis

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	Experim	ental	Contr	ol		Risk Ratio	Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% Cl	M-H, Fixed, 95% Cl
Gao 2015	43	46	41	46	7.7%	1.05 [0.92, 1.19]	- -
Kong 2006	40	47	38	48	7.0%	1.08 [0.89, 1.30]	
Liu 2016	51	59	52	65	9.2%	1.08 [0.92, 1.27]	+
Gao 2014	50	52	45	52	8.4%	1.11 [0.99, 1.25]	
Lv 2015	13	15	11	15	2.1%	1.18 [0.82, 1.70]	
Jin 2013	133	140	106	140	19.8%	1.25 [1.13, 1.39]	
Wang 2012	114	120	90	120	16.8%	1.27 [1.13, 1.42]	
Xing 2018	69	70	53	70	9.9%	1.30 [1.14, 1.49]	
Wen 2014	29	32	22	32	4.1%	1.32 [1.02, 1.71]	
Chen 2017	75	80	48	70	9.6%	1.37 [1.16, 1.62]	
Miao 2014	28	30	20	30	3.7%	1.40 [1.07, 1.83]	
Peng 2017	22	23	9	23	1.7%	2.44 [1.46, 4.10]	
Total (95% CI)		714		711	100.0%	1.24 [1.19, 1.30]	◆
Total events	667		535				
Heterogeneity: Chi ² =	24.90, df =	: 11 (P =	= 0.009);	r = 569	6		
Test for overall effect:	Z = 9.10 (F	• < 0.00	001)				0.5 0.7 1 1.5 2
			,				Favours (experimental) Favours (control)

Figure 2 Forest plot of overall effective rate in safflower yellow group compared with control group.

Table 2 Multivariate	meta-regression	analysis of year	and type of injection

	netvariate meta regress	ion analysis of year and	rtype of injection		
Logor	Coef.	Std.Err.	t	P> t	95% CI
Year	0.0190542	0.0124645	1.53	0.161	-0.0091425, 0.0472508
Туре	0.1216314	0.0704028	1.73	0.118	-0.0376307, 0.2808936
Cons	-38.24614	25.11704	-1.52	0.162	-95.06482, 18.57255

The variant type means safflower yellow injection of the different manufacturer. The proportion of between-study variance explained, R²=0.6586. Coef., regression coefficient; Std.Err., standard error.

	Experim	ental	Conti	ol		Odds Ratio	Odds F	Ratio	
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% Cl	M-H, Fixed	I, 95% CI	
Chen 2017	75	80	48	70	18.2%	6.88 [2.44, 19.38]			
Gao 2014	50	52	43	52	9.4%	5.23 [1.07, 25.54]	-		
Gao 2015	43	46	33	46	12.2%	5.65 [1.49, 21.45]			
Liu 2016	52	59	51	65	32.7%	2.04 [0.76, 5.47]	+		
Lv 2015	13	15	11	15	8.3%	2.36 [0.36, 15.45]			
Miao 2014	28	30	20	30	7.6%	7.00 [1.38, 35.48]		· · · · ·	-
Wen 2014	29	32	22	32	11.7%	4.39 [1.08, 17.89]	-	•	
Total (95% Cl)		314		310	100.0%	4.33 [2.67, 7.05]		•	
Total events	290		228						
Heterogeneity: Chi ² =	: 3.95, df =	6 (P = 0	.68); l² =	0%					
Test for overall effect	: Z = 5.91 (I	P < 0.00	001)				0.01 0.1 1 Favours [experimental]	10 Favours (control)	100

Figure 3 Forest plot of effective rate in safflower yellow group.

conducted through the Chinese 315 Drug Price Inquiry Network (https://www.315jiage.cn). The price of drugs used in the randomized controlled trials was included in the meta-analysis. According to the order of the price per milligram of active ingredient, the sensitivity analysis was conducted with the highest manufacturer retail price and the lowest manufacturer retail price. When calculating the drug cost of the both groups, the combined dose of the most used various drugs or injections in the literature was used as the drug cost of the treatment plan. The results showed that in the safflower yellow group, the treatment measures for calculating the cost were safflower yellow 100 mg

	Experim	ental	Contr	ol		Odds Ratio	Odds Ratio	
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% Cl	M-H, Fixed, 95% Cl	
Jin 2013	133	140	106	140	32.0%	6.09 [2.60, 14.30]		
Kong 2006	40	47	38	48	33.8%	1.50 [0.52, 4.35]		
Peng 2017	22	23	9	23	2.4%	34.22 [3.90, 300.31]		\rightarrow
Wang 2012	114	120	90	120	27.2%	6.33 [2.53, 15.88]		
Xing 2018	69	70	53	70	4.6%	22.13 [2.85, 171.63]		
Total (95% CI)		400		401	100.0%	6.00 [3.71, 9.72]	•	
Total events	378		296					
Heterogeneity: Chi ² =	10.56, df=	= 4 (P =	0.03); l² =	62%			0.01 0.1 1 10	100
Test for overall effect:	Z=7.29 (F	° < 0.00	001)				0.01 0.1 1 10 Favours [experimental] Favours [control]	100

Figure 4 Forest plot of effective rate in safflower injection group.

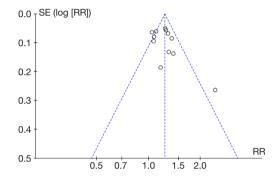


Figure 5 The funnel plot.

Table 3 Inclusion of adverse reactions

Adverse reactions research ID	Slight headache (E/C)	Sickness and vomit (E/C)	Loss of appetite (E/C)	Fever (E/C)	Diarrhea (E/C)	Allergic reaction (E/C)	Incidence rate (E/C)
Peng W, 2017	0/0	1/0	0/0	0/0	0/0	1/3	8.69%/21.73%
Cheng QH, 2017	2/5	3/7	2/6	0/0	0/0	0/0	8.75%/25.71%
Lv XX, 2015	0/1	0/0	1/2	0/1	1/0	0/0	13%/26%
Wen ZP, 2014	1/3	1/2	1/2	0/0	0/0	0/0	9.38%/21.88%
Jin YQ, 2013	0/0	0/2	0/1	0/0	0/0	0/0	0/2.14%

E/C means experimental group/control group.

Table 4 Specific drug price chart

Drug generic name	Cost	Maximum (yuan)	Minimum (yuan)
Safflower yellow injection	36.97 yuan/50 mg ×1 branch	79.9	20.5
Safflower injection	8.95 yuan/5 mL ×1 branch	12.6	2.21
Sanqi Panax notoginseng injection	38.56 yuan/400 mg ×1 branch	20	98
Ozagrel sodium injection	10 yuan/80 mg ×1 branch	11.48	2

Data sources: https://www.315jiage.cn.

Table 5 Results of cost-effectiveness analysis of two treatment options

SPN, Sanqi Panax notoginseng; SYI, Safflower yellow injection; ICER, incremental cost-effectiveness ratio.

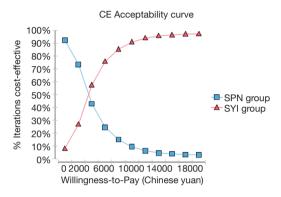


Figure 6 Cost-effective acceptance curve.

combined with ozagrel sodium 80 mg, and the total cost was 1,175.16 RMB (Chinese yuan). See *Table 4* for specific drug prices.

Cost effectiveness analysis

In the case of limited treatment cycle, treatment dose and conventional treatment methods, a total of 12 articles were included in this study, which is the same number of articles included in the meta-analysis, with no article further excluded. According to the proportion of the studies shown in the forest chart in the Meta merger, the treatment efficacy of patients with angina pectoris in each study was weighted to obtain the effectiveness of treatment of cerebral infarction in each group. The results showed that the effective rate of the experiment group was 91.18%, and the effective rate of the Sanqi Panax notoginseng group was 74.83%, which was consistent with the outcome of the meta-analysis, indicating that the weighted calculations had good effectiveness, and could be included in the calculation of pharmacoeconomic evaluation. The results showed that the incremental cost-effectiveness ratio of the safflower yellow group was 3,885.75 RMB with the Sanqi Panax notoginseng group as the control group. According to the recommendations of the World Health Organization on the evaluation of pharmacoeconomics, if ICER < GDP per capita, the increased cost is totally worthwhile; if GDP

per capita < ICER <3 times GDP per capita, the increased cost is acceptable; if ICER > 3 times GDP per capita, the increased cost is not worth it (19)(30). In 2018, China's per capita GDP was 66,006 RMB (32), so the treatment plan in safflower yellow injection group is considered to be completely worthwhile and cost-effective. The detailed results are shown in *Table 5*.

Sensitivity analysis

- (I) Two-factor sensitivity analysis: It was assumed that the treatment efficacy of the two groups fluctuated by 5%; the cost was analyzed using the highest manufacturer retail price and the lowest manufacturer retail price. Assuming that the willingness to pay (WTP) is the GDP per capita in 2018, the whirlwind graph was obtained. The results showed that the parameter that had the greatest impact on the results was the treatment efficiency of the safflower yellow group. When the treatment efficiency of the safflower yellow group changed, the net benefit value of patients varied from 55,999.24 to 6,2018.99 RMB.
- Probability sensitivity analysis: Assumptions were (II)declared that the effective rate was a beta distribution, and the cost was a triangular distribution. After each parameter was put into the TreeAge software, 1,000 second-order Monte Carlo simulations were performed, assuming that the patient's WTP changes ranged from 0 to 20,000 RMB. The cost-effective acceptance curve is shown in Figure 6. It can be seen that when WTP was less than 4,000 RMB, the Sanqi Panax notoginseng (SPN) group had a greater probability of forming a more economical program than the safflower yellow injection (SYI) group; when WTP was higher than 4,000 RMB, the probability of SYI forming a more economical program was greater than 50%. That is, the SYI group is more cost-effective than the SPN group. The results of probability sensitivity analysis were consistent with the results of basic analysis, indicating that the results of basic analysis were robust.

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Discussion

Although Sanqi *Panax notoginseng* and safflower yellow have different mechanisms of action and main components, there are similarities in their pharmacological effects. It has been found through literature review that the relevant studies have been mostly conducted with small sample sizes, and some systematic reviews have been conducted previously to study the treatment of acute cerebral infarction with safflower yellow. No meta-analysis of safflower yellow injection and Sanqi *Panax notoginseng* injection exists in recent literature even though the therapeutic effects of safflower yellow injection and Sanqi *Panax notoginseng* injection on cerebral infarction can be analyzed and compared (8,33). Accordingly, several studies have been used to compare safflower yellow injection in treatment of cerebral infarction through meta-analysis.

In this work, the meta-analysis method was used to compare the effectiveness of injections of safflower yellow and Sanqi Panax notoginseng in the treatment of cerebral infarction, and the Bayesian method was used to compare the cost-effectiveness of the safflower yellow and Sanqi Panax notoginseng. A total of 12 studies were included in this systematic review (13,20-29). The baseline characteristics were compared and not statistically significant in each study. Meta-analysis showed that there was statistical difference in the total effectiveness between the experimental group and the control group when safflower yellow injection and Sanqi Panax notoginseng injection were used to treat acute cerebral infarction. This suggests that the adjuvant treatment using safflower yellow injection or Sanqi Panax notoginseng injection can improve the clinical effects of conventional treatment. From the perspective of treatment effects, using safflower yellow may be a better choice than the Sanqi Panax notoginseng. The evaluation indexes selected in this analysis were consistent, and the use of neurological deficit scores enhanced the rationality and comparability of the evaluation of treatment effects. In terms of safety, only six of the evaluated papers reported the occurrence of adverse reactions, and all of them were minor adverse reactions.

From the perspective of pharmacoeconomics, the ICER is 3,885.75 RMB, that is, the SYI group was more costeffective than SPN group, which is consistent with the result of cost-effectiveness analysis designed by Liu J in 2016. Thus, according to the result information of Sanqi *Panax notoginseng* injection and safflower yellow injection, we recommended that safflower yellow injection should be used to acute cerebral infarction patients priorly to Sanqi

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Panax notoginseng injection in public hospitals.

The limitations of this study include: (I) the small sample size that was finally included in this study somewhat reduces the confidence or widespread applicability of this analysis; (II) the Cochrane score of the final literature reports included in this study is low, and there is lack of sufficient information to judge the research randomization, allocation concealment, blind methods, etc.; (III) the publications included in this study used different doses of safflower yellow pigment, and conventional treatments use different drugs, which may lead to selective bias; (IV) the literature included in this study are all published journal articles, absent of "gray literature", such as special reports, unpublished materials, etc.; (V) all the selected literature reported short-term studies without long-term follow-up monitoring, so the long-term safety and efficacy of safflower vellow treatment could not be evaluated.

Conclusions

In summary, the common use of safflower yellow injection and Sanqi Panax notoginseng injection in the treatment of acute cerebral infarction can improve the overall effectiveness of conventional treatment, and it is safe to use with few adverse reactions. At the same time, in case of high WTP, safflower yellow injection is more costeffective than Sanqi Panax notoginseng injection. Therefore, it is recommended that in the clinical treatment of acute cerebral infarction, safflower yellow pigment injection is used, and safflower yellow pigment injection can be combined with ozagrel sodium for injection within the economically acceptable range of most patients, which is inline with the cost of Sanqi Panax notoginseng. Considering the limited amount of data available for meta-analysis, it is suggested that more clinical trials be undertaken to verify these conclusions with more high-quality data, support further use of these formulations, and evaluate their longterm safety and efficacy profiles.

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Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

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