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Review Article Chinese Medicine Shenfu Injection for Heart Failure: A Systematic Review and Meta-Analysis

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Objective. Heart failure (HF) is a global public health problem. Early literature studies manifested that Shenfu injection (SFI) is one of the most commonly used traditional Chinese patent medicine for HF in China. This article intended to systematically evaluate the efficacy and safety of SFI for HF. *Methods.* An extensive search was performed within 6 English and Chinese electronic database up to November 2011. Ninety-nine randomized controlled trails (RCTs) were collected, irrespective of languages. Two authors extracted data and assessed the trial quality independently. RevMan 5.0.2 was used for data analysis. *Results.* Compared with routine treatment and/or device support, SFI combined with routine treatment and/or device support, SFI combined with routine treatment and/or device support showed better effect on clinical effect rate, mortality, heart rate, NT-proBNP and 6-minute walk distance. Results in ultrasonic cardiography also showed that SFI combined with routine treatment improved heart function of HF patients. There were no significant difference in blood pressure between SFI and routine treatment groups. Adverse events were reported in thirteen trails with thirteen specific symptoms, while no serious adverse effect was reported. *Conclusion.* SFI appear to be effective for treating HF. However, further rigorously designed RCTs are warranted because of insufficient methodological rigor in the majority of included trials.

1. Introduction

Heart failure (HF) is a leading cause of death, hospitalization, and rehospitalization worldwide. Despite advances in the treatment of HF, including use of drugs, devices, and heart transplantation, the condition remains associated with substantial morbidity and mortality [1].

International cooperation research program on cardiovascular disease in Asia showed that, on a total of 15,518 Chinese adults (35–74 years old) survey, the prevalence of HF was 0.9%, 0.7% for the males, and 1.0% for the females [2]. In the United States, HF incidence approaches 10 per 1,000 of the population over 65 years of age [3]. A report from the European Society of Cardiology (ESC) indicated at least 10 million patients with HF in these representing countries with a population of over 900 million. Half of the HF patients will die within 4 years, and more than half of those with severe HF will die within 1 year [4].

At present, the conventional therapeutic approaches in HF management include angiotensin-converting enzyme

(ACE) inhibitors, β -blockers, and diuretics. Although several of them have led to an important effectiveness, HF remains the leading cardiovascular disease with an increasing hospitalization burden and an ongoing drain on health care expenditure [5]. Therefore, it remains necessary to search alternative and complementary treatment, in which Traditional Chinese Medicine takes a good proportion [6].

In TCM theory, pathogenesis of HF is related to deficiency of heart *yang* and heart *qi* and stasis of *blood* and excessive *water* (*fluid*), as well as interaction within these pathological factors. Under physiological conditions, *yang* can promote *water* metabolism, while *qi* can accelerate *blood* circulation, so *yang* and *qi* are the vital elements for human body to maintain life activity. TCM theory holds that patients suffered from HF are in deficiency of heart *yang* and *qi* for a long course, which directly leads to excessive *fluid* retention and *blood* stasis (Figure 1).

Two Chinese herbal medicines, namely, Radix *Ginseng* (ginseng) and Radix *Aconiti Lateralis Preparata* (prepared aconite root), are used in treating HF over 2000 years.

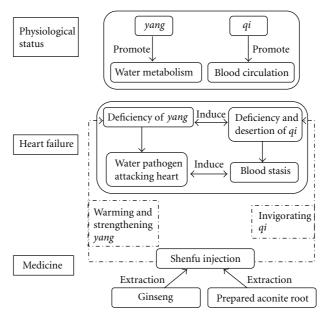


FIGURE 1: TCM theory on heart failure and Shenfu injection.

Ginseng invigorates *qi*, while prepared aconite root can warm and strengthen yang and lead to diuresis. Long-term clinical practice has proved that compatibility of ginseng and prepared aconite root can effectively ameliorate patients' symptom of HF and improve quality of life (Figure 1).

Shenfu injection (SFI) has been used in treating cardiac diseases for a long time in China [7]. The main active components of SFI are extraction of traditional Chinese herbs, namely, ginsenosides and higenamine. Modern pharmacological research shows that ginsenosides can improve ischemic myocardium metabolism, scavenge free radicals, protect myocardial ultrastructure, and reduce Ca²⁺ overload, and higenamine can enhance heart contractility, improve coronary circulation, and decrease the effect of acute myocardial ischemia [8].

Currently, SFI used alone or integrated with routine treatments has been widely accepted as an effective method for the treatment of HF in China. Many clinical studies reported the effectiveness ranging from case reports and case series to controlled observational studies and randomized clinical trials, but the evidence for its effect is not clear. This paper aims to evaluate the beneficial and harmful effects of SFI for treatment of HF in randomized controlled trials.

2. Methods

2.1. Database and Search Strategies. A systematic search was conducted in 5 databases including PubMed (1980–2011), China National Knowledge Infrastructure (1994–2010), VIP Database for Chinese Technical Periodicals (1979–2010), Chinese Biomedical Literature Database (1995–2011), and Cochrane Library (Issue 10, 2011), with the following terms: (Shenfu injection or Shenfu or Shen-fu) AND (heart failure or cardiac dysfunction or cardiac inadequacy or cardiac failure or congestive heart failure). All of those searches ended

before November 2011. And the bibliographies of included trials were searched for thorough references, irrespective of languages.

2.2. Inclusion Criteria. All the randomized controlled trails (RCTs) of SFI compared with routine or conventional treatment (control group) in adult patients with HF were included. RCTs combined SFI with conventional treatment and/or invasive respiratory support (SFI group) compared with conventional treatments and/or invasive respiratory support (control group) were included. Both acute heart failure and chronic heart failure were included. Outcome measures include clinical effect rate, death and adverse events, ultrasonic cardiography, heart rate and blood pressure, and quality of life.

2.3. Data Extraction and Quality Assessment. Two authors (S. Wen-Ting and C. Fa-Feng) extracted the data from the included trials independently, based on the inclusion criteria outlined above. Nonrandomized evaluations, pharmacokinetic studies, animal/laboratory studies, and general reviews were excluded, and duplicated publications reporting the same groups of patients were also excluded (Figure 2).

Extracted data was entered into an electronic database by two authors, S. Wen-Ting and C. Fa-Feng independently. The methodological quality of RCTs was assessed by using criteria from the Cochrane Handbook for Systematic Reviews of Interventions, Version 5.0.1. The quality of trials was categorized into low risk of bias, unclear risk of bias, or high risk of bias according to the risk for each important outcome within included trials, including adequacy of generation of the allocation sequence, allocation concealment, blinding, whether there were incomplete outcome data or selective outcome, or other sources of bias.

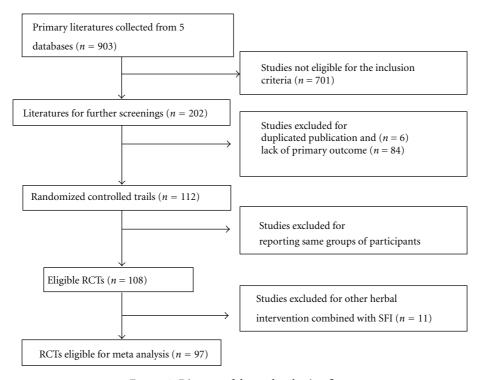


FIGURE 2: Diagram of the study selection flow.

2.4. Data Synthesis. The statistical package (RevMan 5.0.2), which is provided by The Cochrane Collaboration, was used to analyze collected data. Dichotomous data was presented as risk ratio (RR), with 95% confidence intervals (CIs). Continuous outcomes were presented as mean difference (MD), with 95% CI. Analyses were performed by intention-to-treat where possible. Heterogeneity between trials results was tested, and heterogeneity was presented as significant when I^2 is over 50% or P < 0.1. Random effect model was used for the meta-analysis if there was significant heterogeneity, and fixed effect model was used when the heterogeneity was not significant [21]. Publication bias was explored via a funnel-plot analysis.

3. Result

3.1. Search Flow. According to the search strategy, we screened out 903 potentially relevant studies for further identification (Figure 2). By reading titles and abstracts, we excluded 701 studies that were obviously ineligible, including review articles, case reports, animal/experimental studies, and nonrandomized trials. 202 studies with full text papers were retrieved. After the full text reading, 6 studies were excluded because of duplicated publication. 84 studies were excluded due to lack of clinical effect rate which is the primary outcome evaluated in present study. 4 studies were same as previous trials. In 108 RCTs, 11 studies were excluded due to other herbal intervention which was combined with SFI as treatment arm. Thus, 97 RCTs [9–20, 22–108] were included for systematic review.

3.2. Description of Included Trials. Ninety seven RCTs involved a total of 8,202 patients with HF, including 92 trails (7854 patients) of chronic HF and 5 trials (348 patients) of acute HF. The sample size varied from 24 to 248 participants, with an average of 42 patients per group. Since RCTs of HF on children were excluded, patients are adults (ranged from 28 to 89 years old). More males were included than females (52% males and 48% females). Disease duration was reported in 31 trials, ranging from 3 months to 26 years. 49 trials were observed in inpatients, 5 outpatients [22–26], 5 both inpatients and outpatients [27–31], and 39 unclear. All studies were published in Chinese.

Mortality was reported in eleven studies, while the rest of the eighty eight trials did not mention death. Effect rate was assessed in all the trials, based on the improvement of heart function. Ninety one trials used New York Heart Association (NYHA) Classification of Clinical Status, and six trials used Killip's Rating Standards [22, 25, 26, 33–35] for diagnosing HF and rating the patients. Patients in fifty one trials ranged from II to IV, seven trials II to III, twenty one trials III to IV, and five trials IV according to NYHA Classification; patients in five trials ranged from II to IV and one trial IV according to Killip's Standard.

Results of ultrasonic cardiography were reported in 61 trails (5135 patients) with left ventricular ejection fraction (LVEF) as main parameter. Other parameters such as left ventricular diastolic diameter (LVDd), cardiac output (CO), cardiac index (CI), stroke volume (SV), and A peak E-wave velocity ratio (E/A) were reported in 16, 17, 20, 18, and 11 trials, respectively. N-terminal pro-B-type nature tripeptide (NT-proBNP) level in blood was reported in 12 studies of 887 patients, and 6-minute walk distance (6-MWD) was reported

in 8 trials of 630 patients. Heart rate, systolic blood pressure (SBP), and diastolic blood pressure (DBP) were reported in 27, 15, and 13 trials, respectively (Table 1).

3.3. Methodological Quality of Included Trials. According to our predefined quality assessment criteria, all of 97 included trials were evaluated as having unclear risk of bias (Table 2, Figure 3). None of the 97 trials reported sample size calculation. Eleven trials described randomization procedures, nine trials [9-11, 20, 30, 38-41] used a random number table, one drew lots [19], and one trial separated patients by odd and even number of patient ID as a quasirandomization [42]. Only one trial [43] blinded both patients and outcome assessors, and three trials [44-46] blinded patients. None of the trials reported adequate allocation concealment. Five out of ninety seven trials mentioned that followup ranged from 3 months to 12 months after treatment. One trial [47] followed all the patients for 12 months, one trail [38] for 6 month, and the rest [9, 11, 12] for 3 months. However, neither of them used intention to treat method.

3.4. Effect of the Interventions. The primary outcomes were effect rate and mortality. Secondary outcome measures included LVEF, LVDd, SV, CO, CI, HR, systolic blood pressure (SBP), diastolic blood pressure (DBP), NT-proBNP, and 6-MWD.

3.4.1. Primary Outcomes

Effect Rate. All the trials reported clinical effect rate to evaluate the outcome, which was based on NYHA Classification of Clinical Status and Killip's Rating Standards. Killip's Rating Standards were used by six trials with patients of myocardial infarction-induced HF, while other trials used NYHA Classification. Most of trails used three categories to evaluate treatment effect including markedly effective (an improvement of two classes on the classification), effective (an improvement of one class), and ineffective (no improvement, deterioration or death), and others only reported total effect. Total effect rate is the combination of markedly effect rate and effect rate. Trials of myocardial infarction-induced HF and nonmyocardial infarction-induced HF were separated into two subgroups. The meta-analysis showed a total significant difference between SFI and control groups on total effect rate (RR: 1.19, 95% CI [1.17, 1.21]; P < 0.01). And significant difference appeared in both subgroups separately, with RR ratio 1.19 in subgroup of myocardial infarction-induced HF (95% CI [1.16, 1.21]; *P* < 0.01), and 1.46 in the other subgroup (95% CI [1.25, 1.70]; *P* < 0.01) (Figure 4).

Death. Eleven studies reported mortality data, and total death number was 142 out of 978. Two trials [12, 38] assessed the mortality with 3- and 6-month followup, respectively, and other trials reported death at the end of treatment course. Trials were also separated into two subgroups depending on whether HF was induced by myocardial infarction. The result of meta-analysis indicated that SFI can significantly reduce mortality of patients of myocardial

infarction-induced HF (RR: 0.52, 95% CI [0.37, 0.74]; P < 0.01). In the other subgroup, there was no significant difference between mortalities of SFI group and control group (RR: 0.68, 95% CI [0.36, 1.26]; P = 0.22). However, total result of both subgroups showed significant difference (RR: 0.56, 95% CI [0.41, 0.75]; P < 0.01) (Figure 5).

3.4.2. Secondary Outcomes

NT-proBNP. NT-proBNP level is used for screening and diagnosis of acute HF and may be useful to establish prognosis in HF, as it is typically higher in patients with worse outcome [109]. It was reported in 12 studies [20, 22, 38, 45, 49, 52, 54–59] on 887 patients. Consistent with effect rate and other outcomes, NT-proBNP levels of SFI group were significantly lower than control group (WMD: -201.26; 95% CI [-255.27, -147.25], P < 0.01) (Figure 6).

6-*MWD*. Eight trials [47–54] assessed 6-MWD of patients who received SFI or routine treatment. At the end of treatment, eight trails all showed significant increase in walking distance in SFI group, and meta-analysis result was WMD: 14.22; 95% CI [10.31, 18.13], P < 0.01 (Figure 7).

Heart Rate and Blood Pressure. Heart rate and blood pressure were reported in 27 and 15 trials, respectively. Metaanalysis showed that there was statistical significance between SFI group and control group (WMD: 6.31; 95% CI [5.18, 7.44], P < 0.01) (see Supplementary Figure 1 in Supplementary Material available online at doi: 10.1155/2012/713149). However, there was no significant difference between both SBP and DBP in two groups (WMD: -0.07; 95% CI [-0.42, 0.27], P = 0.68) (WMD: -0.37; 95% CI [-0.97, 0.23], P =0.22) (Supplementary Figures 2 and 3).

Results of Ultrasonic Cardiography. LVEF is the ratio of the stroke volume and the left ventricular end-diastolic volume [107]. It is usually used for the assessment of HF and drug efficacy. Sixty-one studies reported the outcomes for LVEF. Meta-analysis showed that SFI group was better than control group in increasing LVEF (WMD: 6.31; 95% CI [5.18, 7.44], P < 0.01) (Supplementary Figure 4).

SV is the volume per stroke by left ventricle, and CO is the volume of blood being pumped by the heart in the time interval of one minute [107]. CI is a vasodynamic parameter that is relating CO to body surface area [107]. All the three parameters indicate left ventricular systolic function, as LVEF does. This paper made meta-analysis of these outcomes, respectively; results showed that SFI group was better than control group in these three parameters: SV (WMD: 7.25; 95% CI [4.60, 9.90], P < 0.01); CO (WMD: 0.67; 95% CI [0.47, 0.87], P < 0.01); CI (WMD: 0.36; 95% CI [0.23, 0.48], P <0.01) (Supplementary Figures 5–7).

E/A ratio is widely accepted as a clinical marker of diastolic HF, and E/A ratio is reduced in diastolic dysfunction [108]. The result of meta-analysis of E/A ratio was WMD: 0.15; 95% CI [0.08, 0.22], P < 0.01, which indicated that SFI better improved diastolic function of heart on HF patients

of including trials.	
ABLE 1: Characters	

Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease duration	Followup (month)
Bao and Yu [61]	Y	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Chen [55]	Unclear	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	Unclear	No
Chen and Liu [14]	Υ	60 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt plus metoprolol 6.25 mg, bid, po	Conventional medicine treatment plus metoprolol 6.25 mg, bid,po	111-11	1–15 y	No
Chen and Li [51]	Υ	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt plus sodium nitroprusside 50 mg, iv.gtt	Conventional medicine treatment plus sodium nitroprusside 50 mg, iv.gtt	IV	Unclear	No
Chen et al. [52]	Υ	15 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	4.5 y on average	No
Chen et al. [56]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	1.5 month–8 y	No
Cui [86]	Unclear	10 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Digoxigenin 0.25 mg	III-II	2-7 y	No
Deng and Tang [15]	Υ	14 d	Conventional medicine treatment plus SFI 20–40 mL, qd, iv:gtt	Conventional medicine treatment	II–IV	Unclear	No
Di [67]	Unclear	Unclear	Conventional medicine treatment plus SFI 40 mL, bid, iv.gtt	Conventional medicine treatment	II–IV	3-17 y	No
Dou [97]	Unclear	10 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	12 ± 1.5 y	No

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Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease duration	Followup (month)
Fan [60]	Y	21 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Metoprolol 12.5 mg, bid, po, +captopril 12.5 mg, tid, po	II-IV	Unclear	No
Fan et al. [101]	Unclear	14 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Geng et al. [27]	Both	12 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	0.5–9 y	No
Gu et al. [69]	Υ	14 d	Conventional medicine treatment plus SFI 100 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	1.5–12 y	No
Guo et al. [49]	Unclear	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	Unclear	No
Guo et al. [23]	Z	7 d	Conventional medicine treatment plus SF1 20 mL, iv + 50 mL, qd, iv.gtt plus non invasive positive pressure ventilation	Conventional medicine treatment plus non invasive positive pressure ventilation	Unclear	Unclear	No
Guo et al. [102]	Υ	7 d	Conventional medicine treatment plus SFI 40–60 mL, qd, iv.gtt plus invasive respiratory support	Conventional medicine treatment plus invasive respiratory support	IV	Unclear	No
Han and Li [36]	Υ	15 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	AI-III	$4.54 \pm 2.1\mathrm{y}$	No
He [70]	Unclear	14 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	1-14 y	No

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TABLE	

Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease duration	Followup (month)
He [98]	Unclear	7–20 d/ 10–30 d	Conventional medicine treatment plus SFI 20–40 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	3-16у	No
Hong [44]	Unclear	14 d	Conventional medicine treatment plus SFI 80 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	Unclear	No
Hou and Hong [17]	Unclear	7 d	Conventional medicine treatment plus SFI 60–100 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Huang [13]	Unclear	7 d	Conventional medicine treatment plus SFI 20 mL iv + 40 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	Unclear	No
Huang [53]	Υ	14 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Huang et al. [24]	Z	Unclear	Conventional medicine treatment plus SFI 50 mL, qd, ivgtt plus sodium nitroprusside 50 mg, ivgtt	Conventional medicine treatment plus sodium nitroprusside 50 mg, iv.gtt	Unclear	Unclear	No
Jia and Yang [71]	Υ	20 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Jian and Chen [88]	Unclear	14 d	Conventional medicine treatment plus SFI 60–80 mL, bid, iv.gtt	Conventional medicine treatment	AI-III	6.5 y on average	No
Jiang [62]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Jin and Guo [95]	Y	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No

Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease duration	Followup (month)
Ju [37]	Unclear	14 d	Conventional medicine treatment plus SFI 30 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Lei and Li [92]	Υ	7–10 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Lei et al. [12]	Y	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	11–IV	1–18 <i>y</i>	ω
Li et al. [9]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	11–IV	1-20 y	ω
Li et al. [72]	Υ	15 d	Conventional medicine treatment plus SFI 30 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	5-26 y	No
Li [96]	Unclear	15 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	Unclear	No
Li et al. [73]	Unclear	15 d	Conventional medicine treatment plus SFI 100 mL, qd, iv.gtt plus sodium nitroprusside 50 mg	Conventional medicine treatment plus sodium nitroprusside 50 mg	IV	l-25 y	Q
Li [93]	Unclear	10 d	Conventional medicine treatment plus SFI 1 mL/kg body weight, qd, iv.gtt	Conventional medicine treatment	11–IV	Unclear	No
Liu [75]	Y	21 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	11–1V	Unclear	No

Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease H duration	Followup (month)
Liu and Sun [18]	Unclear	7 d	Conventional medicine treatment plus SFI 100 mL, qd, iv.gtt	Conventional medicine treatment	Unclear	Unclear	No
Liu and Chan [50]	Unclear	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Liu et al. [20]	Υ	28 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	9 month–14 y	No
Liu [74]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Liu et al. [94]	Unclear	14 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Lv [57]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	Unclear	No
Luo et al. [76]	Х	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment plus sodium nitroprusside	VI-III	Unclear	No
Luo et al. [38]	Unclear	10 d/m 6 months	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	>3 months	Q
Ma et al. [48]	Unclear	20 d	Conventional medicine treatment plus SFI 30–40 mL, qd, iv.gtt	Conventional medicine treatment	111-11	Unclear	No
Ma and Huang [99]	Υ	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No

TABLE 1: Continued.

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Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease duration	Followup (month)
Ma [77]	Unclear	15 d	Conventional medicine treatment plus SFI 20 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Pan et al. [89]	Y	14 d	Conventional medicine treatment plus SFI 100 mL, qd, iv.gtt	Conventional medicine treatment plus dobutamine hydrochloride 40 ng, qd, iv.gtt	III-IV	2.5 month–11 <i>y</i>	No
Qiu [103]	Y	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Ru [46]	Unclear	10 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	Unclear	No
Shang [78]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Song [106]	Υ	15 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	111-111	Unclear	No
Song et al. [10]	Y	15 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Su [90]	Y	14 d	Conventional medicine treatment plus SFI 100 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Tàn et al. [58]	Unclear	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Tian and Gong [16]	Υ	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	2-20 y	No

			TABLE 1: Continued.	ed.			
Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease duration	Followup (month)
Tian [80]	Y	15 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	NI-III	>7 months	No
Tu and Yang [63]	Υ	14 d	Conventional medicine treatment plus SFI 80 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Tu et al. [32]	Unclear	14 d	Conventional medicine treatment plus SFI 100 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
G. L. Wang and J. Wang [104]	Υ	15 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	2.5–16 y	No
Wang [100]	Υ	14 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	Unclear	$22.3 \pm 4.8\mathrm{y}$	No
Wang [39]	Unclear	15 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Wang [28]	Both	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	0.6-7 y	No
Wang and Ye [87]	Υ	10 d	Conventional medicine treatment plus SFI 40–100 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	14.2 y mean	No
Wang et al. [81]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	IV	3-10 y	No
Wu and Duan [45]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	III-II	Unclear	No
Wu and Wang [64]	Y	14 d	Conventional medicine treatment plus SFI 100 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No

			TABLE 1. CONTINUECO.	.u.			
Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease duration	Followup (month)
Wu et al. [40]	Unclear	10 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Yang and Wu [82]	Y	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	AI-III	Unclear	No
Yang et al. [54]	Y	15 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	5.1 y	No
Yao and Lu [65]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Yin [83]	Υ	14 d	Conventional medicine treatment plus SFI 40 mL, qd, iv	Conventional medicine treatment	VI-II	0.5–12 y	No
Yu et al. [84]	Unclear	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Yu [66]	Unclear	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	11–1V	Unclear	No
Zhan and Yang [47]	Unclear	20 d	Conventional medicine treatment plus SFI 40 mL, qd, iv.gtt plus metoprolol. 25 mg–75 mg, bid po	Conventional medicine treatment plus metoprolol. 25 mg–75 mg, bid, po	111-11	Unclear	12
Zhang [79]	Y	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	3–15 y	No
Zhang et al. [85] 	Y	20 d	Conventional medicine treatment plus SFI 30 mL, qd, iv.gtt	Conventional medicine treatment	VI-II	Unclear	No

Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease duration	Followup (month)
Zhang [42]	Y	21 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	III-II	Unclear	No
Zhang and Pan [30]	Both	14 d	Conventional medicine treatment plus SFI 40–60 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	2-16 y	No
Zhang [29]	Both	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	VI-III	Unclear	No
Zhang [43]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Zhao et al. [11]	Υ	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment plus isoket 10 mg,qd,iv.gtt	II-IV	1–20 y	ŝ
Zhao [91]	Υ	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	Unclear	Unclear	No
Zhou [59]	Υ	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	IV	3–15 y	No
Zhou [19]	Υ	10 d	Conventional medicine treatment plus SFI 80 mL, qd, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Zhou et al. [31]	Both	14 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Zhu and Ma [105]	Υ	15 d	Conventional medicine treatment plus SFI 50 mL, qd, ivgtt	Conventional medicine treatment	VI-III	Unclear	No

TABLE 1: Continued.

			TABLE 1. COULING	acu.			
Author Name	Inpatient (Y/N)	Course	Experiment group	Control group	NYHA classification	Disease duration	Followup (month)
Zi and Li [41]	Y	14 d	Conventional medicine treatment plus SFI 40–100 mL, qd, iv or iv.gtt	Conventional medicine treatment plus dobutamine hydrochloride 50–100 mg	II–IV	Unclear	No
Guo et al. [22]	Z	14 d	Conventional medicine treatment plus SFI 60–100 mL, bid, iv.gtt	Conventional medicine treatment	II-IV	Unclear	No
Mo and Zhao [25]	Z	7 d	Conventional medicine treatment plus SFI 60–100 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	1–7 d	No
Song and Zhang [33]	Υ	10 d	Conventional medicine treatment plus SFI 40–60 mL, qd, iv.gtt	Conventional medicine treatment plus dobutamine hydrochloride 40 mg	II-IV	Unclear	No
Zeng et al. [34]	Υ	7 d	Conventional medicine treatment plus SFI 50 mL, qd, iv.gtt	Conventional medicine treatment	IV	Unclear	No
Zeng [26]	Z	10 d	Conventional medicine treatment plus SFI 60–100 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	1-72 h	No
Zhang [35]	Y	14 d	Conventional medicine treatment plus SFI 60 mL, qd, iv.gtt	Conventional medicine treatment	II–IV	Unclear	No
Conventional medicine treatmand β -blockers.	ent includes sitting	up position, suppl	Conventional medicine treatment includes sitting up position, supplemental oxygen, vasodilator such as nitroglycerine, diuretics such as furosemide, and cardiotonic agents such as lanatoside C, ACE inhibitors, and β -blockers.	/cerine, diuretics such as furosemi	ide, and cardiotonic agent:	s such as lanatoside C,	ACE inhibitors,

Author Name	Sequence generation	Allocation concealment	Blinding	Incomplete outcome data	Selective outcome reporting	Other source of bias	Risk of bias
Bao and Yu [61]	Unclear	Unclear	Z	N	z	Unclear	Unclear
Chen [55]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Chen and Liu [14]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Chen and Li [51]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Chen et al. [52]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Chen et al. [56]	Unclear	Unclear	Z	Υ	Z	Unclear	Unclear
Cui [86]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Deng and Tang [15]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Di [67]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Dou [97]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Fan [60]	Unclear	Unclear	Z	Ν	Z	Unclear	Unclear
Fan et al. [101]	Unclear	Unclear	Ν	Ν	Z	Unclear	Unclear
Gao et al. [68]	Unclear	Unclear	Ν	Ν	Z	Unclear	Unclear
Geng et al. [27]	Unclear	Unclear	Ν	N	Z	Unclear	Unclear
Gu et al. [69]	Unclear	Unclear	Ν	N	Z	Unclear	Unclear
Guo et al. [49]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Guo et al. [23]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Guo et al. [102]	Unclear	Unclear	Z	Υ	Z	Unclear	Unclear
Han and Li [36]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
He [70]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
He [98]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Hong [44]	Unclear	Unclear	Single-blind	Z	Z	Unclear	Unclear
Hou and Hong [17]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Huang [13]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Huang [53]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Huang et al. [24]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Jia and Yang [71]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Jian and Chen [88]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Jiang [62]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Jin and Guo [95]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Ju [37]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Lei and Li [92]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Lei et al. [12]	Unclear	Unclear	Z	Υ	Z	Unclear	Unclear
Li et al. [9]	Random number table	Unclear	Z	Z	Z	Unclear	Unclear
Li et al. [72]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
ון:[מע]							

	Sequence generation	Allocation concealment	Blinding	Incomplete outcome data	Selective outcome reporting	Other source of bias	Risk of bias
Li et al. [73]	Unclear	Unclear	N	N	Z	Unclear	Unclear
Li [93]	Unclear	Unclear	Z	Ν	Z	Unclear	Unclear
Liu [75]	Unclear	Unclear	Z	Ν	Z	Unclear	Unclear
Liu and Sun [18]	Unclear	Unclear	N	N	Z	Unclear	Unclear
Liu and Chan [50]	Unclear	Unclear	Z	Z	Ν	Unclear	Unclear
Liu et al. [20]	Random	Unclear	Z	Υ	Z	Unclear	Unclear
I i [74]	Tinulosi tauto	Thelear	Ν	Λ	Ν	Hnclear	Thelear
Liu et al. [94]	Unclear	Unclear	ΖZ	- Z	άZ	Unclear	Unclear
Lv [57]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Luo et al. [76]	Unclear	Unclear	Z	Z	N	Unclear	Unclear
Luo et al. [38]	Random number table	Unclear	Z	Υ	Ν	Unclear	Unclear
Ma et al. [48]	Unclear	Unclear	N	Z	Ν	Unclear	Unclear
Ma and Huang [99]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Ma [77]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Pan et al. [89]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Qiu [103]	Unclear	Unclear	Z	Ν	Z	Unclear	Unclear
Ru [46]	Unclear	Unclear	Single-blind	N	Z	Unclear	Unclear
Shang [78]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Song [106]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Song et al. [10]	Random number table	Unclear	Z	Z	Z	Unclear	Unclear
Su [90]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Tan et al. [58]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Tian and Gong [16]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Tian [80]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Tu and Yang [63]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Tu et al. [32]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
G. L. Wang and J. Wang [104]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Wang [100]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Wang [39]	Random number table	Unclear	Z	Z	Z	Unclear	Unclear
Wang [28]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Wang and Ye [87]	Unclear	Unclear	Z	Υ	Z	Unclear	Unclear
Wang et al. [81]	Unclear	Unclear	Z	N	Z	Unclear	Unclear

Author Name	Sequence generation	Allocation concealment	Blinding	Incomplete outcome data	Selective outcome reporting	Other source of bias	Risk of bias
Wu and Duan [45]	Unclear	Unclear	Single-blind	Y	Z	Unclear	Unclear
Wu and Wang [64]	Unclear	Unclear	Z	Ν	Z	Unclear	Unclear
Wu et al. [40]	Random number table	Unclear	Z	Z	Z	Unclear	Unclear
Yang and Wu [82]	Unclear	Unclear	Ζ	Ν	Ν	Unclear	Unclear
Yang et al. [54]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Yao and Lu [65]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Yin [83]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Yu et al. [84]	Unclear	Unclear	Ν	Z	Ν	Unclear	Unclear
Yu [66]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Zhan and Yang [47]	Unclear	Unclear	Z	Υ	Z	Unclear	Unclear
Zhang [79]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Zhang et al. [85]	Unclear	Unclear	Ν	Z	Ν	Unclear	Unclear
Zhang [42]	odd and even number of ID	Unclear	Z	Z	Z	Unclear	Unclear
Zhang and Pan [30]	Unclear	Unclear	Z	Z	Ζ	Unclear	Unclear
Zhang [29]	Random number table	Unclear	Z	Z	Z	Unclear	Unclear
Zhang [43]	Unclear	Unclear	Double-blind	Z	Z	Unclear	Unclear
Zhao et al. [11]	Random number table	Unclear	Z	Z	Z	Unclear	Unclear
Zhao [91]	Unclear	Unclear	Z	N	Z	Unclear	Unclear
Zhou [59]	Unclear	Unclear	Ν	Z	Z	Unclear	Unclear
Zhou [19]	Drew lots	Unclear	Ν	Z	Z	Unclear	Unclear
Zhou et al. [31]	Unclear	Unclear	Ν	Z	Z	Unclear	Unclear
Zhu and Ma [105]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Zi and Li [41]	Random number table	Unclear	Z	Z	Z	Unclear	Unclear
Guo et al. [22]	Unclear	Unclear	Ν	Υ	Ν	Unclear	Unclear
Mo and Zhao [25]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Song and Zhang [33]	Unclear	Unclear	Z	Υ	Z	Unclear	Unclear
Zeng et al. [34]	Unclear	Unclear	Z	Z	Ν	Unclear	Unclear
Zeng [26]	Unclear	Unclear	Z	Z	Z	Unclear	Unclear
Zhang [35]	Unclear	Unclear	z	Y	Z	Unclear	Unclear

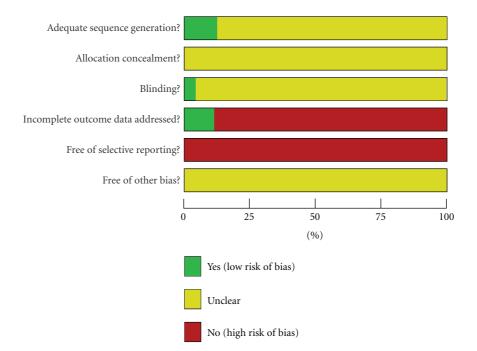


FIGURE 3: Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.

	TABLE 5: Adverse events	
Symptom	Reported trails	Cases reported
Dry mouth	4 [10, 16, 17, 60]	14
Fullness of the head	4 [9–12]	10
Dryness heat	2 [10, 13]	7
Insomnia	1 [13]	3
Dysphoria	1 [14]	2
Skin itching	1 [15]	1
Tachycardia	1 [16]	1
Feverish dysphoria	2 [17, 18]	5
Flushing of face and tidal fever	1 [19]	8
Dizziness due to low blood pressure	1 [20]	1
Gastrointestinal discomfort	1 [20]	1
Palpitation	1 [18]	2

TABLE 3. Adverse events

than conventional medicine treatment did (Supplementary Figure 8).

LVDd is the end-diastolic dimension of the left ventricle. There was no statistical significance between SFI combined with conventional medicine treatment and conventional medicine treatment groups (WMD: -1.59; 95% CI [-5.29, 2.12], P = 0.40) (Supplementary Figure 9).

3.4.3. Quality of Life. None of the trials reported quality of life.

3.5. Publication Bias. Funnel plots based on the data of effect rate were elaborated in Figure 8. The figure was asymmetrical, which indicated that potential publication bias might influence the results of this paper. Although we conducted comprehensive searches and tried to avoid bias, since all trials were published in Chinese, we could not exclude potential publication bias.

3.6. Adverse Effect. Thirty seven out of ninety seven trials mentioned the adverse effect except in sixty-two trials which was unclear. Thirteen trials [9–20, 60] reported the following thirteen specific symptoms of side effects including dry mouth, dryness heat, fullness of the head, insomnia, dysphoria, skin itching, tachycardia, feverish dysphoria, flushing of face, tidal fever, dizziness due to low blood pressure, gastro-intestinal discomfort, and palpitation. Among these side effects, dry mouth and fullness of the head were reported in 4 trails with 14 and 10 cases, respectively. These symptoms were regarded to be mild and recovered spontaneously after SFI withdrawal. Twenty four trials reported that no side effects were observed in the SFI group (Table 3).

The above side effects might be related to higenamine, which is the active ingredient of prepared aconite root. In TCM books and papers, prepared aconite root is frequently mentioned with adverse effects as dry mouth, dryness heat, fullness of the head, and dysphoria due to its strong effect of strengthening *yang*.

4. Discussion

In many years, western medicine has made tremendous progress and has become the dominating medical treatment worldwide. However, it has been increasingly recognized that

Study or Subgroup Events Total Events Total Weight IV, fixed, 95% CI V, fixed, 95% CI V, fixed, 95% CI V, 2011 V, 2017		Shenfu i	njection	Contr	ol		Risk ratio	Risk ratio
Bao G. H. 2011 27 30 20 39 0.4% 175 [126, 2,44] Chem 14. 2007 31 33 52 33 0.4% 175 [126, 2,44] Chem 34. 2009 37 40 32 40 1.2% 116 0.57, 1.38 Chem 34. 2009 37 40 32 40 1.2% 116 0.57, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 37 41 112 40.4% 124 1.02, 1.38 Chem 34. 2009 38 43 30 122 50 0.34% 135 (0.38, 1.26) Chem 34. 2009 38 43 31 32 30 0.7% 132 (0.38, 1.26) Chem 34. 2006 31 33 32 40 7% 124 (0.38, 1.26) Chem 34. 2006 31 33 32 40 7% 124 (0.38, 1.26) Chem 34. 2006 31 33 32 21 30 0.7% 132 (0.38, 1.26) Chem 34. 2006 31 33 32 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2006 31 33 32 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2006 31 33 32 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2008 31 33 32 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 38 43 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 38 43 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 38 43 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 38 43 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 38 43 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 38 43 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 38 43 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 38 43 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 38 43 22 30 0.7% 124 (0.38, 1.26) Chem 34. 2009 34 43 24 44 2.3% 114 (0.38, 1.26) Chem 34. 2009 34 43 24 44 2.3% 114 (0.38, 1.26) Chem 34. 2007 44 44 44 2.3% 144 (0.38, 1.26) Chem 34. 2007 44 44 44 2.3% 144 (0.38, 1.26) Chem 34. 2007 44 44 44 2.3% 144 (0.38, 1.26) Chem 34. 2007 44 44 44 2.3% 147 (0.38, 1.26) Chem 34. 2007 44 44 44 2.3% 146 (1.25, 1.17) Chem 34. 2007 44 44 44 44 2.3% 1						-	IV, fixed, 95% C	I IV, fixed, 95% CI
Chen J, H. 2007 31 33 32 33 0.5% 133 102 124 3 Chen X. L. 2009 32 33 33 32 27 0.9% 135 0.3% 134 0.27.13 Chen X. L. 2009 32 33 02 27 0.9% 135 0.3% 134 0.7% 135 0.2	Bao G. H. 2011	27	30	20	39		1.75 [1.26, 2.44]]
Chen X. L. 2009 Gui Z. J. 2000 Gui Z. 2	Chen J. H. 2007	31	35	22	33	0.5%	1.33 1.02, 1.74	
Cui L. 2000, 79 9 90 6 11 90 128 13 11, 1, 1, 12 Dou J. 2008, 77 47 14 14 22 04.98 13 11, 1, 1, 12 Dou J. 2008, 77 47 14 14 12 04.98 13 11, 1, 1, 12 Dou J. 2008, 77 47 14 14 12 04.98 13 Cuo J. 2009, 75 46 14 14 12 04.98 13 Cuo J. 2009, 75 46 14 14 12 04.98 144 10.05, 198 Cuo J. 2008, 73 41 14 12 20 4.98 1.56 11.08, 128 Cuo J. 2008, 73 41 14 12 20 4.98 1.56 11.08, 128 Cuo J. 2008, 73 41 14 12 20 4.98 1.56 11.08, 128 Cuo J. 2008, 73 43 13 32 80 0.78 1.56 11.08, 128 Cuo J. 2008, 73 43 15 32 80 0.78 1.02 10.81, 129 Cuo J. 2008, 74 48 42 73 01.28 1.15 0.27, 136 Han W.F. 1999 46 48 42 73 01.28 1.15 0.27, 136 Han W.F. 1999 46 48 42 73 01.28 1.13 0.027, 136 Han W.F. 1999 46 48 42 73 01.28 1.13 10.05, 168 Han W.F. 1999 46 48 42 73 01.28 1.13 10.05, 168 Han W.F. 1999 46 48 42 73 00 1.28 1.13 10.05, 168 Han W.F. 1999 46 48 42 73 00 1.28 1.13 10.05, 168 Han W.F. 1999 46 48 42 73 00 1.23 1.106, 1586 Han W.F. 1999 46 48 42 33 00 0.478 1.23 10.05, 168 Han W.F. 1999 46 48 42 33 00 0.478 1.23 10.05, 168 Han W.F. 1999 46 48 42 33 00 0.478 1.23 10.05, 168 Han W.F. 1999 46 48 42 34 0.478 1.23 10.05, 158 Han W.F. 1999 46 48 42 44 48 2.48 1.12 0.98, 153 Han W.G. 2006 40 44 44 48 42 42 48 1.100, 129, 157 Han W.G. 2008 40 46 46 198 122 1.106, 1586 Huang W.O. 2008 40 46 52 40 43 0.49 1.23 10.05, 114 Huang W.O. 2008 40 44 52 34 0.148 1.23 1.100, 1146 Huang W.O. 2008 41 43 23 33 00 .88 1.120 0.98, 153 Han W.G. 2000 45 50 44 48 44 48 2.288 1.12 0.98, 154 Huang W.O. 2008 44 48 44 48 2.288 1.11 0.05, 114 Huang W.O. 2008 44 48 44 48 2.288 1.11 0.05, 114 Huang W.O. 2008 44 48 41 48 2.288 1.12 0.089, 169 Hu H. 2009 45 50 34 64 53 40 1.13 1.00, 125 Hu H. 2009 45 50 34 64 33 40 1.18 1.20 1.14 11 Huang W.O. 2008 54 42 33 54 1.148 1.29 1.100, 1141 Huang W.O. 2008 54 42 33 54 1.148 1.29 1.100, 123 1.14 Huang W.O. 2008 54 42 33 54 1.148 1.29 1.100, 123 1.14 Huang W.O. 2008 54 42 33 54 1.148 1.29 1.100, 114 11 Huang W.O. 2008 54 44 53 54 00 43 4.120 0.98, 1.38 1.100, 1.20 1.141 Huang W.O. 2008 54 44 53 54 00 4	Chen X. L. 2009	32	35	26	35	0.8%	1.23 0.99, 1.53	
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	Chen Z. G. 2009 Cui Z. J. 2000	79	90	61	90	1.5%	1.15[0.93, 1.4] 1.3[1.1, 1.52]	. T—
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	Deng X. Y. 2011 Di S. T. 2010					1.5% 4.2%	1.09 [0.93, 1.28] 1.20 [1.09, 1.32]	
$ \begin{bmatrix} \operatorname{Pan} S, M, 2010 \\ \operatorname{Cons} X, Y, 1000 \\ \operatorname{Cons} X, $	Dou J. 2008	37	41		22	0.4%	1.42 [1.02, 1.98]	
Gu X, 50,2005 Gu D, L2006 313 Gu D, L2006 313 Gu D, L2006 313 Gu D, L2006 313 Gu D, L2006 313 Gu D, L2006 313 Gu D, L2006 313 414 Horg M, L2000 17 18 15 18 14 12 10 12 12 12 12 12 12 12 12 12 12	Fan S. M. 2010	36	40	30	40	0.9%	1.2 [0.98, 1.48]	, 1
Guo L. 2008 Guo Y. E. 2009 Han W. F. 1999 Han W. L. 2000 Han W. L. 2008 Han W. Han W. H	Geng X. Y. 2006	30	40	12	20	0.2%	1.25 [0.84, 1.86]	j <u> </u>
Han W. F. 1999 46 48 25 30 1.3% 1.13 0.57, 1.36 Hong M. L. 2000 35 36 24 30 0.3% 1.13 0.51, 1.68 Hong M. L. 2004 36 24 26 39 0.6% 1.33 1.05, 1.68 Hong M. L. 2004 36 24 26 26 90 6.4% 1.31 0.25, 1.68 Hung H. 1009 36 38 27 38 0.6% 1.33 1.05, 1.68 Hung W. C. 2004 36 48 26 39 0.6% 1.33 1.05, 1.68 Hung W. C. 2005 37 62 56 62 3.3% 1.02 0.91, 1.14 Hung W. C. 2005 37 62 56 62 3.3% 1.02 0.91, 1.14 Hung W. C. 2006 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2006 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 50 39 760 2.35% 1.35 1.09, 1.52 H. 2.00 70 88 41 54 1.49% 1.22 0.98, 1.52 H. 2.00 70 88 41 54 1.49% 1.22 0.97, 1.34 Hung W. C. 2009 42 7 50 39 760 2.35% 1.35 1.01, 1.81 H. 2.00 70 82 47 12 0.04% 1.25 0.97% 1.35 1.01, 1.81 H. 2.00 70 82 47 12 0.04% 1.25 0.97% 1.35 1.01, 1.81 H. 2.00 70 82 47 12 0.04% 1.12 0.97, 1.37 Hu X. 2.009 36 44 0 32 40 1.2% 1.04 0.46, 1.25 1.14 Hung X. 2.009 36 44 0 32 40 1.15% 1.04 0.86, 1.25 1.14 Hung X. 2.009 36 44 0 32 4.33% 1.17 0.99, 1.39 Hu X. 2.009 36 44 0 32 4.39% 1.13 0.97, 1.37 Hu X. 2.009 36 44 0 32 4.39% 1.13 0.97, 1.37 Hu X. 2.000 31 64 62 33 40 1.19% 1.16 0.96, 1.14 Hu X. 2.000 31 64 62 33 40 1.19% 1.16 0.96, 1.14 Hu X. 2.000 31 64 62 33 40 1.19% 1.04 0.86, 1.25 1.14 Hu X. 2.000 31 82 22 41 40 0.6% 1.35 1.01, 1.81 Hu X. 2.000 31 82 22 41 40 0.6% 1.35 1.01, 1.81 Hu X. 2.000 31 82 22 41 40 0.6% 1.36 1.03 Hu X. 2.001 37 85 46 0 40 52 0.33% 1.10 0.98, 1.36 1.14 Hu X. 2.001 37 84 69 40 42 1.14 0.98, 1.36 1.14 Hu X. 2.001 37 84 69 40 42 1.14 0.98, 1.36 1.14 Hu X. 2.001 37 84 60 40 52 0.33% 1.10 0.98, 1.36 1.14 Hu X. 2.001 37 84 60 40 52 0.33% 1.10 0.98, 1.36 1.14 Hu X. 2.001 37 84 60 40 52 0.33% 1.10 0.98, 1.36 1.14 Hu X. 2.001 37 84 40 22 34 100 0.98 1.1	Guo J. 2008	26	31	23	28	0.7%	1.02 [0.81, 1.29]	
Han W. F. 1999 46 48 25 30 1.3% 1.13 0.57, 1.36 Hong M. L. 2000 35 36 24 30 0.3% 1.13 0.51, 1.68 Hong M. L. 2004 36 24 26 39 0.6% 1.33 1.05, 1.68 Hong M. L. 2004 36 24 26 26 90 6.4% 1.31 0.25, 1.68 Hung H. 1009 36 38 27 38 0.6% 1.33 1.05, 1.68 Hung W. C. 2004 36 48 26 39 0.6% 1.33 1.05, 1.68 Hung W. C. 2005 37 62 56 62 3.3% 1.02 0.91, 1.14 Hung W. C. 2005 37 62 56 62 3.3% 1.02 0.91, 1.14 Hung W. C. 2006 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2006 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 03 34 0.15% 1.14 0.97, 1.34 Hung W. C. 2009 47 65 50 39 760 2.35% 1.35 1.09, 1.52 H. 2.00 70 88 41 54 1.49% 1.22 0.98, 1.52 H. 2.00 70 88 41 54 1.49% 1.22 0.97, 1.34 Hung W. C. 2009 42 7 50 39 760 2.35% 1.35 1.01, 1.81 H. 2.00 70 82 47 12 0.04% 1.25 0.97% 1.35 1.01, 1.81 H. 2.00 70 82 47 12 0.04% 1.25 0.97% 1.35 1.01, 1.81 H. 2.00 70 82 47 12 0.04% 1.12 0.97, 1.37 Hu X. 2.009 36 44 0 32 40 1.2% 1.04 0.46, 1.25 1.14 Hung X. 2.009 36 44 0 32 40 1.15% 1.04 0.86, 1.25 1.14 Hung X. 2.009 36 44 0 32 4.33% 1.17 0.99, 1.39 Hu X. 2.009 36 44 0 32 4.39% 1.13 0.97, 1.37 Hu X. 2.009 36 44 0 32 4.39% 1.13 0.97, 1.37 Hu X. 2.000 31 64 62 33 40 1.19% 1.16 0.96, 1.14 Hu X. 2.000 31 64 62 33 40 1.19% 1.16 0.96, 1.14 Hu X. 2.000 31 64 62 33 40 1.19% 1.04 0.86, 1.25 1.14 Hu X. 2.000 31 82 22 41 40 0.6% 1.35 1.01, 1.81 Hu X. 2.000 31 82 22 41 40 0.6% 1.35 1.01, 1.81 Hu X. 2.000 31 82 22 41 40 0.6% 1.36 1.03 Hu X. 2.001 37 85 46 0 40 52 0.33% 1.10 0.98, 1.36 1.14 Hu X. 2.001 37 84 69 40 42 1.14 0.98, 1.36 1.14 Hu X. 2.001 37 84 69 40 42 1.14 0.98, 1.36 1.14 Hu X. 2.001 37 84 60 40 52 0.33% 1.10 0.98, 1.36 1.14 Hu X. 2.001 37 84 60 40 52 0.33% 1.10 0.98, 1.36 1.14 Hu X. 2.001 37 84 60 40 52 0.33% 1.10 0.98, 1.36 1.14 Hu X. 2.001 37 84 40 22 34 100 0.98 1.1	Guo J.J. 2006 Guo Y. F. 2009	33 71	36 76	19 42	26 76	$0.6\% \\ 0.9\%$	1.25 [0.97, 1.62] 1.69 [1.37, 2.09]	
Hong M. L. 2000 Hong M. L. 2000 Hung E. 2009 Hung E. 2009 Hung E. 2009 Hung W. 2008 Hung K. 2	Han W. F. 1999	46	48	25	30	1.3%	1.15 0.97, 1.36	
Hou S. L. 2004 42 48 26 39 0.0% 1.31 1.03, 1.68 Hum g. L. 1999 36 38 228 30 0.9% 1.23 1.05, 1.86 Hum g. V. 02 2008 57 62 56 62 3.2% 1.06 0.91, 1.14 Hum g. V. 02 2008 57 62 56 62 3.2% 1.06 0.91, 1.14 Hum g. V. 02 2007 68 84 43 61 1% 1.22 1.04, 1.46 Hum g. V. 02 2007 68 84 43 61 1% 1.22 1.04, 1.46 Hum g. V. 02 2007 68 84 43 60 1% 1.22 1.04, 1.46 Hum g. V. 2007 68 84 43 60 1% 1.22 1.04, 1.46 Hum g. V. 2009 57 56 33 40 1.5% 1.14 0.93, 1.16 Hum g. V. 2009 53 54 44 31 48 2.2% 1.12 0.88, 1.28 Hu Y. S. 2007 68 84 63 60 53.4% 1.04 0.93, 1.16 Hu Y. S. 2007 64 48 41 48 2.2% 1.12 0.88, 1.28 Hu Y. S. 2007 64 48 41 48 2.2% 1.12 0.88, 1.28 Hu Y. S. 2007 64 48 41 48 2.2% 1.12 0.88, 1.28 Hu Y. S. 2007 64 48 41 48 2.2% 1.12 0.88, 1.28 Hu Y. S. 2007 64 55 9 39 50 1.3% 1.16 0.86, 1.28 Hu Y. S. 2007 64 55 9 39 50 1.3% 1.16 0.86, 1.28 Hu Y. J. 2009 63 64 63 32 44 01 1.40 (0.68, 1.28 Hu Y. J. 2006 68 72 64 43 34 01 1.49 (0.68, 1.28 Hu Y. J. 2006 71 55 03 93 50 1.3% 1.16 0.86, 1.21 Hu X. J. 2009 36 64 23 40 1.14% 1.00 (0.86, 1.21 Hu Y. J. 2006 31 32 29 32 448 1.107 0.94, 1.21 Hu Y. J. 2006 31 42 22 37 11 20 0.68, 1.28 Hu Y. J. 2006 31 42 21 41 0.20 Hu Y. J. 2006 31 42 21 41 0.20 41 42 0.5% 1.28 Hu Y. J. 2006 31 32 42 1.28 Hu Y. J. 2007 145 50 0.38 50 1.14% 1.16 0.96, 1.21 Hu Y. J. 2008 34 60 40 52 1.33% 1.17 (0.94, 1.21 Hu Y. J. 2008 34 66 40 52 1.33% 1.17 (0.94, 1.21 Hu Y. J. 2008 34 67 32 44 11 48 1.16 0.18, 1.28 Hu H. J. 2001 27 32 117 30 0.4% 1.38 1.06, 1.48 Hu H. J. 2001 27 32 117 30 0.4% 1.38 1.01, 1.47 Hu H. J. 2001 27 32 41 73 00 0.4% 1.28 1.16 (0.95, 1.34 Hu H. J. 2001 27 32 41 73 00 0.4% 1.28 1.16 (0.95, 1.38 Hu H. J. 2001 33 44 23 44 0.0% 1.28 1.16 (0.18, 1.28] Hu H. J. 2001 33 44 23 44 0.0% 1.28 1.16 (0.18, 1.28] Hu H. J. 2009 34 47 72 93 66 1.19% 1.14 (0.95, 1.38 Hu H. J. 39 Hu H. J. 2011 35 62 Hu H. J. 39 Hu H. J. 2011 35 62 Hu H. J. 39 Hu H. J. 104 Hu H. J. 39 Hu H. J. 2011 35 64 60 38 35 0 1.2% 1.13 Hu H. J. 39 Hu H. J. 2011 36 88 22 50 0.7% 1.1	He X. J. 2006	55	60	44	60	1.3%	1.25 [1.05, 1.48]	
Hung L, 2009 36 38 22 38 0.9% 1.33 1.09, 1.65 Han C, 2005 2008 47 46 26 46 03 32% 10.20 31, 134 Han C, 2007 68 80 40 60 196 122 104, 1.56 Han C, 2007 68 80 40 60 196 122 104, 1.56 Han C, 2007 68 80 40 60 196 122 104, 1.56 Han C, 2007 68 14 14 0.33, 116 Lei W, 2000 53 34 42 21 44 0.3% 1.51 109, 2.15 Lei W, 2000 53 34 42 21 44 0.3% 1.51 109, 2.15 Lei W, 2000 53 34 42 34 134 0.23% 1.16 Lei W, 2000 53 35 44 34 14 14% 1.22 104, 1.53 Li C, 2.007 68 38 44 34 14 14% 1.22 104, 1.54 Lei W, 2000 54 38 44 34 144% 1.22 104, 1.54 Lei W, 2000 54 38 44 34 144% 1.22 104, 1.54 Li C, 2.007 46 48 133% 1.16 0.86, 1.23 Li C, H. 2004 63 66 52 64 2.3% 1.17 103, 1.34 Li Z, H. 2004 63 66 52 64 2.3% 1.17 103, 1.34 Li Z, H. 2004 63 66 52 64 2.3% 1.17 103, 1.34 Li Z, H. 2004 63 66 52 64 1.35 Li Z, H. 2007 45 50 39 50 1.3% 1.16 0.86, 1.25 Li W X, 2009 36 44 33 40 1.2% 1.04 0.86, 1.25 Li W X, 2009 36 44 32 29 32 2.44% 10.70 0.94, 1.21 Lu X, 2006 51 42 23 40 1.2% 1.06 0.66, 1.4 Li X, 2.000 52 91 21 21 40 0.66 1.25 Li W X, 2009 34 45 03 38 50 1.1% 1.16 0.96, 1.4 Li U X, 1.2008 51 42 23 42 41 100, 103, 1.37 Li U X, 2009 34 47 50 338 50 1.1% 1.16 0.96, 1.4 Lu X, 2.000 52 91 22 43 80 0.4% 1.38 1.02, 1.87 Lu X, 2.000 54 46 21 24 42 10.60, 1.29 Li W X, 2000 53 43 72 29 36 1.1% 1.16 0.35, 1.37 Li U X, 2000 73 45 70 46 53 1.53% 1.17 0.99, 1.39 Ha S, B. 2011 47 50 46 53 1.53% 1.17 0.99, 1.39 Ha S, B. 2011 47 50 46 53 1.53% 1.17 0.99, 1.39 Ha S, B. 2011 47 50 46 53 1.53% 1.17 0.99, 1.39 Ha S, B. 2011 45 50 22 26 1.1% 1.16 0.88, 1.28 Ha S, B. 2011 47 50 46 63 0.40 2.3 1.3% 1.44 Ha S, B. 2011 33 42 83 41 42% 1.18 (1.53, 1.38 Ha S, B. 2011 33 42 83 41 42% 1.18 (1.53, 1.38 Ha S, B. 2011 33 42 83 41 42% 1.14 (0.58, 1.38] Ha L, 2000 73 43 72 29 36 1.1% 1.04 (0.38, 1.28] Ha M, 2000 73 43 72 29 36 1.1% 1.46 (0.58, 1.28] Ha M, 2000 73 44 72 20 0.7% 1.27 (1.01, 1.61 Ha M, W, 2010 73 74 02 24 30 0.7% 1.27 (1.03, 1.61 Ha M, W, 2000 74 43 72 73 73 9.84% 1.19 (1.06, 1.51) Ha M, W, 2000 74 48 30 42 90 0.4% 1.23 (0.88, 1.28]	Hou X. L. 2004	42	48	26	39	0.6%	1.31 [1.03, 1.68]	
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	Huang T. 2009	36	38	27	38	0.8%	1.33 [1.07, 1.66]	
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	Huang W. Q. 2008 Jia Q. 2005			28 56		0.7% 3.2%	1.24 [0.99, 1.57] 1.02 [0.91, 1.14]	
$ \begin{array}{c} \mu_{1,3,2} 2009, \\ \mu_{1,3,2} 2009, \\ \mu_{1,3,2} 2009, \\ \mu_{1,3,2} 2000, \\ \mu_{1,3,2} 2000, \\ \mu_{1,2,2} 2000, \\ \mu_{2,3,3} 20, \\ \mu_{2,3,3} $	lian Y. P. 2002		64	47	64	1.4%	1.23 1.04, 1.46	
Lei W. G. 2003 33 42 21 41 0.3% 1.53 1.09.2.15 Li D. 2010 54 58 41 44% 1.23 1.04.1.453 Li D. 2007 63 66 51 24 0.3% 1.23 1.04.1.453 Li D. 2007 63 66 51 24 2.3% 1.17 1.03.1.34 Li D. 2009 24 28 14 20 0.4% 1.22 0.88 1.69 Li U. 2009 24 28 14 20 0.4% 1.22 0.89 1.69 Li U. 2009 24 28 14 20 0.4% 1.22 0.89 1.69 Li U. 2009 24 22 35 01 1.9% 1.04 0.86 1.25 Li U. Y. 2009 46 74 63 78 2.4% 1.14 1.00 1.29 Li U. X. 2009 46 74 63 78 2.4% 1.14 1.00 1.29 Li U. Y. 2005 40 42 33 40 1.2% 1.04 0.86 1.25 Li U. Y. 2005 40 42 33 40 1.2% 1.09 0.91, 1.3] Lu O. X. 2009 43 32 29 32 2.4% 1.07 (0.94, 1.2] Lu O. X. 2009 43 32 29 32 2.4% 1.07 (0.94, 1.2] Lu O. X. 2009 44 33 32 29 32 2.4% 1.07 (0.94, 1.2] Lu O. X. 2009 44 33 32 29 32 2.4% 1.07 (0.94, 1.2] Lu O. X. 2009 44 33 32 29 32 2.4% 1.07 (0.94, 1.2] Lu O. X. 2009 44 33 32 29 30 0.4% 1.83 1.02, 1.57 J. Ma J. J. 2003 54 60 40 50 3.3% 1.02 (0.92, 1.14) Ma J. J. 2003 54 61 62 50 3.3% 1.02 (0.92, 1.14) Ma J. J. 2003 85 46 71 82 2.13% 1.11 (0.92, 1.15) Ma S. B. 2011 37 56 60 50 60 2.23% 1.12 (0.98, 1.28) J. 2011 56 60 50 60 2.23% 1.11 (0.92, 1.13) Ma S. B. 2011 33 34 28 34 1.4% 1.18 [1.129, 1.81] Song S. Q. 1099 39 45 33 42 19% 1.11 (0.93, 1.134] J. 2003 18 226 23 1.16 0.138 [1.02, 1.81] J. 2013 34 28 34 1.4% 1.18 [0.138, 1.28] J. 2014 49 58 25 38 0.6% 1.28 [1.166 [0.38, 1.28] J. 2014 49 58 25 1.18 (0.93, 1.44] J. 2015 34 44 27 34 0.6% 1.28 [1.06, 1.48] J. 2014 49 58 25 1.18 (0.16, 1.88, 1.28] J. 30 1.16 (0.88, 1.28] J. 30 1.16 (0.88, 1.28] J. 30 1.13 1.6 (0.98, 1.28] J. 31 1.05 1.16 J. 30 34 44 27 34 0.6% 1.23 [1.01, 1.51] J. 40 37 400 36 40 1.30 (0.38, 1.28] J. 41 (0.93, 1.35] J. 41 (0.93, 1.36] J. 41 (0.93, 1.36] J. 41 (0.93, 1.36] J. 41 (0.93, 1.36] J. 42 J. 200 37 40 32 40 0.3% 1.32 [1.05, 1.6] J. 42 0.98 37 40 32 40 0.3% 1.32 [1.05, 1.6] J. 20 0.83 51 44 27 34 0.6% 1.23 [1.01, 1.51] J. 40 Ma R. W. 2009 37 40 32 40 0.3% 0.138 [1.07, 1.38] J. 41 (0.93, 1.38 0.77, 1.37] J. 41 (0.93, 1.38 0.77, 1.37] J. 41 (0.93, 1.38 0.77, 1.37] J. 41 (0.42, 27	Jin X. P. 2007	28	30	23	30	0.8%	1.22 [0.98, 1.52]	
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$ \begin{bmatrix} v G_{2} 010 & 29 & 31 & 21 & 30 & 0.6\% & 1.34 & [1.04, 1.72] \\ Ma H, W. 2005 & 40 & 42 & 42 & 0.5\% & 1.67 & 1.27, 2.18 \\ Ma J, 2008 & 54 & 60 & 40 & 52 & 1.3\% & 1.17 & 0.99, 1.39 \\ Ma S, B. 2011 & 47 & 50 & 46 & 50 & 3.3\% & 1.02 & 0.92, 1.14 \\ Pan M, 1, 2003 & 28 & 32 & 19 & 30 & 0.4\% & 1.38 & [1.02, 1.87] \\ Pan M, 1, 2001 & 23 & 84 & 41 & 44.6 & 1.18 & [1.09, 1.31] \\ Shang Y. 2011 & 36 & 60 & 50 & 60 & 2.2\% & 1.12 & [0.98, 1.28] \\ Song J, I, 2011 & 33 & 34 & 28 & 34 & 1.4\% & 1.18 & [1.39] \\ Song S, Q. 1999 & 39 & 45 & 33 & 42 & 19\% & 1.11 & [0.91, 1.34] \\ Song S, Q. 1999 & 39 & 45 & 33 & 42 & 19\% & [1.10, 91, 1.34] \\ Tan L, 12011 & 36 & 38 & 22 & 30 & 0.7\% & [1.29] & [1.03, 1.62] \\ Tian I, 2009 & 34 & 37 & 29 & 36 & [1.10, 95, 1.38] \\ Tian I, 2000 & 13 & 61 & 10 & 16 & 0.2\% & [1.5] & [0.83, 2.03] \\ Tian Y, 2010 & 13 & 61 & 10 & 16 & 0.2\% & [1.5] & [0.83, 1.48] \\ Wang W, G. 2006 & 28 & 31 & 24 & 31 & 0.6\% & [1.86] & [0.85, 1.38] \\ Wang W, C. 2006 & 28 & 31 & 24 & 31 & 0.6\% & [1.86] & [0.85, 1.38] \\ Wang W, M. 2009 & 54 & 58 & 43 & 58 & 1.4\% & [1.26] & [1.06, 1.48] \\ Wang W, M. 2009 & 54 & 58 & 43 & 58 & 1.4\% & [1.26] & [1.06, 1.48] \\ Wang W, W. 2008 & 37 & 40 & 30 & 40 & 1\% & [1.23] & [1.03, 1.61] \\ Wang Y. M. 2008 & 37 & 40 & 32 & 40 & 0.5\% & [1.2] & [0.93, 1.55] \\ Wu H, Y. 2010 & 34 & 40 & 24 & 34 & 0.6\% & [1.2] & [0.93, 1.48] \\ Wang Y. M. 2008 & 37 & 40 & 32 & 40 & 0.2\% & [1.27] & [0.38, 1.48] \\ Wang Y. M. 2009 & 24 & 30 & 40 & 100 & (0.96 & 1.22] & [1.93, 1.55] \\ Wu H, Y. 2010 & 37 & 40 & 22 & 30 & 0.7\% & [1.27] & [1.0, 1.161] \\ Yang Y. 2008 & 37 & 40 & 32 & 40 & 0.2\% & [1.20, 93, 1.48] \\ Tan H, 2008 & 54 & 60 & 38 & 50 & 1.2\% & [1.20, 1.48] \\ Tan H, 2008 & 54 & 60 & 38 & 50 & 1.2\% & [1.30, 0.3\% & [1.38] & [1.30, 0.5\% & [1.27] & [1.30, 1.61] \\ Tan H, 2008 & 54 & 50 & 0.5\% & [1.26] & [1.41] & [0.93, 1.48] \\ Tan H, 2008 & 54 & 50 & 0.5\% & [1.26] & [1.41] & [0.93, 1.48] \\ Tan H, 2008 & 11 & [1.44] & 12 & [1.26, 1.44] & [1.46] & [1.44] & [1.45] \\ Tan H, 2008 & 53 & 52 & 55 & 54 & 65 & [1.29\% & [1.48] & [0.$	Liu Y. J. 2005 Luo S. P. 2008					$\frac{1.1\%}{2.4\%}$	1.22 [1.01, 1.47] 1.07 [0.94, 1.21]	
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $						1.1%	1.16[0.96, 1.4] 1 34 [1 04, 1 72]	
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	Ma H. W. 2005	40	42	24	42	0.5%	1.67 [1.27, 2.18]	
Qiu W. W. 2010 Ru H. G. 2001 Shang Y. 2011 56 60 50 60 228 112 1092, 1.3] Tan Song S. Q. 1999 39 45 33 42 144 1.18 1.18 1.1992, 1.3] Tan Song S. Q. 1999 39 45 33 42 144 1.18 1.1991, 1.34] Tan L, 2013 Tan L, 2010 13 16 10 16 0.2% 1.2 1.03, 1.62 1.3 1.04, 2.03 1.3 1.26, 2.16 1.3 1.26, 2.16 1.3 1.26, 2.16 1.26, 1.26 1.26, 1.27 1.21, 1.20 1.21, 1.21, 1.21 1.21, 1	Ma S. B. 2011	47	50	46	50	3.3%	1.02 [0.92, 1.14]	· -
Snang Y. 2011 56 60 50 60 2.5% 1.12 [0.98, 1.28] Song Y. 2019 33 34 53 40 14% 1.18 [1.1.39] Song S. Q. 1999 39 44 53 40 19% 1.17 [0.99, 1.24] Final L. 2001 16 22 41 10 0.7% 1.24 [0.95, 1.38] Tian L. N. 2010 13 16 10 16 0.25% 1.3 [0.83, 2.03] Tu Y. P. 2010 49 58 25 38 0.6% 1.26 [1.26, 2.16] Wang Q. 2009 18 20 15 20 0.5% 1.2 [0.90, 1.61] Wang Y. 2009 54 58 43 58 1.4% 0.168 [0.85, 1.38] Wang X. M. 2009 54 58 43 58 1.4% 0.168 [0.85, 1.38] Wang X. M. 2009 54 58 43 58 1.4% 0.168 [0.85, 1.38] Wang Y. 2009 54 58 43 58 1.4% 0.168 [0.85, 1.38] Wang Y. 2009 54 58 43 58 1.4% 0.126 [1.06, 1.48] Wang Y. 2009 54 58 43 22 30 0.7% 1.27 [1.01, 1.61] Wang Y. 2008 37 40 32 40 1.2% 0.1% 1.55] Wu Y. B. 2008 41 44 27 44 0.6% 1.52 [1.19, 1.95] Wu Y. B. 2008 37 40 32 40 1.2% 0.1% 0.1% 1.51] Yang Y. 2008 28 30 22 30 0.7% 1.27 [1.01, 1.61] Yang Y. 2008 28 30 22 30 0.7% 1.27 [1.01, 1.61] Yang Y. 2008 27 30 21 30 0.6% 1.28 [0.99, 1.67] Yin H. 2008 50 55 62 11 30 0.6% 1.28 [0.99, 1.67] Yin H. 2008 50 55 62 11 30 0.6% 1.28 [0.99, 1.67] Yin H. 2008 54 60 38 40 1.2% 0.188 [0.98, 1.3] Yang Y. 2004 37 40 32 40 1.2% 1.16 [0.97, 1.38] Yang Y. 2008 37 40 32 40 1.2% 1.16 [0.97, 1.38] Yang Y. 2008 36 40 30 40 1.39 [1.05, 1.6] Yin H. 2008 54 60 38 40 1.2% 0.188 [0.99, 1.4] Yu G. Y. 2004 37 59 27 37 0.9% 1.32 [1.05, 1.6] Zhang Y. 2004 37 59 27 37 0.9% 1.32 [1.05, 1.6] Zhang Y. 2003 11 12 8 12 0.2% 1.38 [0.89, 1.2] Zhang Y. 2010 37 46 22 30 0.7% 1.27 [1.01, 1.61] Zhang Y. 2010 53 56 40 56 1.2% 0.188 [0.89, 1.2] Zhang Y. 2001 32 80 22 30 0.7% 1.27 [1.01, 1.61] Zhang Y. 2010 53 56 40 56 1.2% 0.138 [0.89, 1.2] Zhang Y. 2003 28 30 22 30 0.7% 1.27 [1.01, 1.61] Zhang Y. 2010 53 56 40 56 1.2% 0.138 [0.89, 1.2] Zhang Y. 2003 28 30 22 30 0.7% 1.27 [1.01, 1.61] Zhang Y. 2003 28 30 22 30 0.7% 1.27 [1.01, 1.61] Zhang Y. 2003 28 30 22 30 0.7% 1.27 [1.01, 1.61] Zhang Y. 2001 45 50 45 05 0.19% 1.32 [1.07, 1.64] Zhang Y. 2001 21 22 [1.28, 1.48 [0.97, 1.37] Zhang Y. 2001 21 23 [1.20, 1.41 [0.7, 2.3] Zhang Y. 2001 21 23 [1.60, 0.	Oiu W W 2010	77	85	67	85	2.3%	1.15 [1.01, 1.31]]
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	Ru H. G. 2001 Shang Y. 2011					1.3% 2.2%	$1.1 \ [0.92, 1.3]$ $1.12 \ [0.98, 1.28]$	
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	Song J. J. 2011 Song S. O. 1999				34 42	$^{1.4\%}_{1\%}$	1.18[1, 1.39] 1.1[0.91, 1.34]	·
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$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	Tian J. 2009	34	37	29	36	1.1%	1.14 [0.95, 1.38]	<u>+</u>
$\begin{array}{l c c c c c c c c c c c c c c c c c c c$	Tu Q. Y. 2003	52	62	31	61	0.2%	1.65 [1.26, 2.16]]
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Wang G. L. 2011		50	22	26	1.1%	1.06 0.88, 1.28	
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Wang W. G. 2006					0.6%	1 08 [0 85 1 38]	
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Wang W. M. 2009 Wang X M 2009				24 58	0.8% 1.4%	1.1 [0.89, 1.36]	
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Tang L. 2006 28 30 22 30 0.7% 1.27 [1.01, 1.61] Yang Z. Y. 2008 37 40 32 40 1.29 [0.97, 1.38] Yao J. 2007 27 30 21 30 0.6% 1.29 [0.97, 1.38] Yao J. 2008 50 56 21 30 0.6% 1.29 [0.97, 1.38] Yu J. Y. 2010 37 40 28 40 0.8% 1.32 [1.05, 1.64] Yu J. Y. 2010 37 40 28 40 0.8% 1.32 [1.06, 1.65] Zhang L. 2009 54 60 38 50 1.2% 1.18 [0.98, 1.48] Zhang H. 2009 54 60 38 50 1.2% 1.18 [0.98, 1.48] Zhang W. X. 2003 11 12 8 12 0.2% 1.38 [0.98, 1.48] Zhang Y. 2003 11 12 8 12 0.2% 1.38 [0.98, 1.48] Zhang Y. 2003 22 30 0.7% 1.27 [1.01, 1.61] Zhan Y. 2003 28 30 22 30 0.7% 1.22 [1.05, 1.44] Zhang Y. 2003 17 19 13 18 0.4% 1.22 [1.05, 1.44] Zhang Y. 2010 53 56 40 56 1.2% 1.32 [1.07, 1.64] Zhang Y. 2010 55 55 40 56 1.2% 1.13 [0.98, 1.3] Zhou J. 2011 52 56 46 56 1.9% 1.13 [0.98, 1.3] Zhou Z. 7. 2010 28 30 22 30 0.7% 1.27 [1.01, 1.61] Zhao X. 2010 55 55 40 50 1.9% 1.25 [1.08, 1.44] Zi Y. 1999 18 20 13 20 0.3% 1.38 [0.97, 1.97] Heterogeneity: $\chi^2 = 104.42, df = 90 (P = 0.14); I^2 = 14(%)$ Test for overall effect: Z = 16.86 (P < 0.00001) 1.1.2 myocardial infarction induced heart failure Guo Q. 2009 22 35 17 35 0.2% 1.29 [0.85, 1.98] Mo C. R. 2002 24 36 16 38 0.2% 1.58 [1.02, 2.45] Song Q. 2011 21 23 14 22 0.3% 1.43 [1.02, 2.02] Lag Y. 2011 40 54 29 56 0.4% 1.43 [1.02, 2.45] Song Q. 2011 21 23 14 22 0.3% 1.43 [1.02, 2.45] Song Q. 2011 24 36 16 38 0.2% 1.58 [1.02, 2.45] Song Y. L. 2011 40 54 29 56 0.4% 1.43 [1.07, 7.23] Zhang H. 2.011 24 36 16 38 0.2% 1.58 [1.02, 2.45] Song Y. L. 2011 40 54 29 56 0.4% 1.43 [1.02, 2.45] Song Y. L. 2011 40 54 29 56 0.4% 1.44 [1.25, 1.7] Total events 153 104 Heterogeneity: $\chi^2 = 0.62, df = 5 (P = 0.99); I^2 = 0(%)$ Test for overall effect: Z = 4.74 (P < 0.00001) Total (95% CI) 4259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 3831 $= 2874$ Heterogeneity: $\chi^2 = 17.32 (P < 0.00001)$ Total events $\chi^2 = 17.32 (P < 0.00001)$ Total events $\chi^2 = 17.32 (P < 0.00001)$	Wu Y. B. 2008	41	44	27	44	0.6%	1.52 [1.19, 1.95]	į <u> </u>
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Zhang Y. 2011 45 50 34 50 0.9% 1.32 [1.07, 1.64] Zhang Z. M. 2003 28 30 22 30 0.7% 1.27 [1.01, 1.61] Zhao H. 2009 70 78 59 80 1.7% 1.22 [1.05, 1.41] Zhao X. X. 2010 53 56 40 56 1.2% 1.32 [1.05, 1.41] Zhou J. S. 2011 52 56 46 56 1.9% 1.13 [0.98, 1.3] Zhou Z. T. 2010 28 30 22 30 0.7% 1.27 [1.01, 1.61] Zhu C. Z. 2010 55 55 40 50 1.9% 1.23 [1.08, 1.44] Zi Y. 1999 18 20 13 20 0.3% 1.38 [0.97, 1.97] Subtotal (95% CI) 4047 3731 98.4% 1.19 [1.16, 1.21] Total events $3678 - 2770$ Heterogeneity: $\chi^2 = 104.42$, $df = 90$ ($P = 0.14$); $l^2 = 14(\%)$ Test for overall effect: Z = 1.686 ($P < 0.00001$) 1.1.2 myocardial infarction induced heart failure Guo Q. 2009 22 35 17 35 0.2% 1.59 [0.85, 1.98] Mo C. R. 2002 24 36 16 38 0.2% 1.58 [1.02, 2.45] Song Q. 2001 21 23 14 22 0.3% 1.43 [1.02, 2.02] Zhang Y. 2001 20 22 8 12 23 0.2% 1.51 [0.97, 2.33] Zeng Y. 2001 21 23 14 22 0.3% 1.43 [1.06, 1.93] Zhang H.X. 2011 40 54 29 56 0.4% 1.43 [1.06, 1.93] Zhang H.X. 2011 24 36 16 38 0.2% 1.58 [1.02, 2.45] Subtotal (95% CI) 212 212 1.6% 1.46 [1.25, 1.7] Total events 153 104 Heterogeneity: $\chi^2 = 0.62$, $df = 5$ ($P = 0.99$); $l^2 = 0(\%)$ Test for overall effect: Z = 4.74 ($P < 0.00001$) Total (95% CI) 4259 3943 100.0% 1.19 [1.17, 1.21] Heterogeneity: $\chi^2 = 11.32$ ($P < 0.0001$) Total (95% CI) 212 1.2% 1.2% 1.2% 1.2% 1.2% 1.2% 1.4% Heterogeneity: $\chi^2 = 1.732$ ($P < 0.0001$) Total events 2 3831 Heterogeneity: $\chi^2 = 1.732$ ($P < 0.0001$) Total events 2 2 1.1% 40 54 29 56 0.4% 1.19 [1.17, 1.21] Heterogeneity: $\chi^2 = 1.732$ ($P < 0.0001$) Total events 2 153 1004 Heterogeneity: $\chi^2 = 1.732$ ($P < 0.0001$) Total events 2 173 ($P < 0.0001$) Total events 2 1732 ($P < 0.0001$) Total events 2 1732 ($P < 0.00001$) Total e	Zhang L. 2005	29	30	22	26	1.2%	1.14 [0.96, 1.36]	
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Zhang Y. 2011	45	50	34	50	0.9%	1.32 [1.07, 1.64]	
Zhao X, X. 2010 53 56 40 56 1.2% 1.32 [1.11, 1.58] Zhou G, 2010 17 19 13 18 0.4% 1.24 [0.89, 1.72] Zhou J, S. 2011 52 56 46 56 1.9% 1.13 [0.98, 1.3] Zhou Z, T. 2010 28 30 22 30 0.7% 1.27 [1.01, 1.61] Zhu C, Z. 2010 55 55 40 50 1.9% 1.25 [1.08, 1.44] Zhu C, Z. 2010 55 55 40 50 1.9% 1.25 [1.08, 1.44] Zhu C, Z. 2010 55 55 40 70 1.27 [1.01, 1.61] Zhu C, Z. 2010 55 55 40 70 1.29 [0.85, 1.98] Mu C, Z. 2010 4047 3731 98.4% 1.19 [1.16, 1.21] Total events 3678 2770 Heterogeneity: $\chi^2 = 104.42, df = 90 (P = 0.14); l^2 = 14(\%)$ Test for overall effect: $Z = 16.86 (P < 0.0001)$ 1.1.2 myocardial infarction induced heart failure Guo Q, 2009 22 35 17 35 0.2% 1.29 [0.85, 1.98] Mo C, R. 2002 24 36 16 38 0.2% 1.58 [1.02, 2.45] Song Q, 2001 21 23 14 22 0.3% 1.43 [1.06, 1.93] Zhang H, X. 2011 40 54 29 56 0.4% 1.43 [1.06, 1.93] Zhang H, X. 2011 24 36 16 38 0.2% 1.58 [1.02, 2.45] Subtotal (95% CI) 212 212 1.6% 1.46 [1.25, 1.7] Total events 153 104 Heterogeneity: $\chi^2 = 0.62, df = 5 (P = 0.99); l^2 = 0(\%)$ Test for overall effect: $Z = 4.74 (P < 0.00001)$ Total events 3831 2874 Heterogeneity: $\chi^2 = 11.71, df = 96 (P = 0.13); l^2 = 14\%$ Test for overall effect: $Z = 17.32 (P < 0.00001)$ Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2831 2874 Heterogeneity: $\chi^2 = 11.71, df = 96 (P = 0.03); l^2 = 14\%$	Zhang Z. M. 2003 Zhao H. 2009	70	78	59	80	0.7% 1.7%	1.27 [1.01, 1.61] 1.22 [1.05, 1.41]	
Zhou J. S. 2011 52 56 46 56 1.9% 1.13 [0.98, 1.3] Zhou Z. T. 2010 28 30 22 30 0.7% 1.27 [1.01, 1.61] Zhu C. Z. 2010 55 55 40 50 1.9% 1.25 [1.08, 1.44] Zi Y. 1999 18 20 13 20 0.3% 1.38 [0.97, 1.97] Subtotal (95% CI) 4047 3731 98.4% 1.19 [1.16, 1.21] Total events 3678 2770 Heterogeneity: $\chi^2 = 104.42, df = 90 (P = 0.14); l^2 = 14(\%)$ Test for overall effect: Z = 16.86 (P < 0.00001) 1.1.2 myocardial infarction induced heart failure Guo Q. 2009 22 35 17 35 0.2% 1.29 [0.85, 1.98] Mo C. R. 2002 24 36 16 38 0.2% 1.58 [1.02, 2.45] Song Q. 2009 22 28 12 23 0.2% 1.51 [0.97, 2.33] Zeng Y. 2009 22 28 12 23 0.2% 1.51 [0.97, 2.33] Zeng Y. L. 2011 40 54 29 56 0.4% 1.43 [1.06, 1.93] Zhang H.X. 2011 24 36 16 38 0.2% 1.58 [1.02, 2.45] Subtotal (95% CI) 212 212 1.6% 1.46 [1.25, 1.7] Total events 153 104 Heterogeneity: $\chi^2 = 0.62, df = 5(P = 0.99); l^2 = 0(\%)$ Test for overall effect: Z = 4.74 (P < 0.00001) Total (95% CI) 24259 3943 100.0% 1.19 [1.17, 1.21] Total events 3831 4259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 3831 4259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 2 17.32 (P < 0.00001) Total (95% CI) 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 150 14259 3943 100.0% 1.19 [1.17, 1.21] Total events 150 14259 394	Zhao X. X. 2010	53 17	56	40	56	1.2% 0.4%	1.32 [1.11, 1.58] 1.24 [0.89, 1.72]	
Subtotal (95% CI) 4047 3/31 98.4% 1.19 [1.16, 1.21] total events 3678 2770 Heterogeneity: $\chi^2 = 104.42$, $df = 90$ ($P = 0.14$); $l^2 = 14(\%)$ Test for overall effect: $Z = 16.86$ ($P < 0.00001$) 1.1.2 myocardial infarction induced heart failure Guo Q, 2009 22 35 17 35 0.2% 1.29 [0.85, 1.98] Mo C, R. 2002 24 36 16 38 0.2% 1.58 [1.02, 2.45] Song Q, 2001 21 23 14 22 0.3% 1.43 [1.02, 2.02] Zeng Y, L 2011 40 54 29 56 0.4% 1.43 [1.02, 2.45] Subtotal (95% CI) 212 212 1.6% 1.46 [1.25, 1.7] Total events 153 104 Heterogeneity: $\chi^2 = 0.62$, $df = 5$ ($P = 0.99$); $l^2 = 0(\%)$ Test for overall effect: $Z = 4.74$ ($P < 0.00001$) Total (95% CI) 4259 3943 100.0% 1.19 [1.17, 1.21] Heterogeneity: $\chi^2 = 11.32$ ($P < 0.0001$) Total (95% CI) 5831 0.00% 1.19 [1.17, 1.21] Heterogeneity: $\chi^2 = 11.73$, $df = 96$ ($P = 0.13$); $l^2 = 14\%$	Zhou I \$ 2011	52	56	46	56	1.9%	1.13 [0.98, 1.3]	. —
Subtotal (95% CI) 4047 3/31 98.4% 1.19 [1.16, 1.21] total events 3678 2770 Heterogeneity: $\chi^2 = 104.42$, $df = 90$ ($P = 0.14$); $l^2 = 14(\%)$ Test for overall effect: $Z = 16.86$ ($P < 0.00001$) 1.1.2 myocardial infarction induced heart failure Guo Q, 2009 22 35 17 35 0.2% 1.29 [0.85, 1.98] Mo C, R. 2002 24 36 16 38 0.2% 1.58 [1.02, 2.45] Song Q, 2001 21 23 14 22 0.3% 1.43 [1.02, 2.02] Zeng Y, L 2011 40 54 29 56 0.4% 1.43 [1.02, 2.45] Subtotal (95% CI) 212 212 1.6% 1.46 [1.25, 1.7] Total events 153 104 Heterogeneity: $\chi^2 = 0.62$, $df = 5$ ($P = 0.99$); $l^2 = 0(\%)$ Test for overall effect: $Z = 4.74$ ($P < 0.00001$) Total (95% CI) 4259 3943 100.0% 1.19 [1.17, 1.21] Heterogeneity: $\chi^2 = 11.32$ ($P < 0.0001$) Total (95% CI) 5831 0.00% 1.19 [1.17, 1.21] Heterogeneity: $\chi^2 = 11.73$, $df = 96$ ($P = 0.13$); $l^2 = 14\%$	Zhu C. Z. 2010	55	55	40	50	1.9%	1.25 [1.08, 1.44	
Heterogeneity: $\chi^2 = 104.42$, $df = 90$ ($P = 0.14$); $I^2 = 14(\%)$ Test for overall effect: $Z = 16.86$ ($P < 0.00001$) 1.1.2 myocardial infarction induced heart failure Guo Q. 2009 22 35 17 35 0.2% 1.29 [0.85, 1.98] Mo C. R. 2002 24 36 16 38 0.2% 1.43 [1.02, 2.45] Zeng Y. 2009 22 28 12 23 0.2% 1.43 [1.02, 2.02] Zeng Y. 2009 22 28 12 23 0.2% 1.43 [1.02, 2.02] Zeng Y. 2011 24 36 16 38 0.2% 1.43 [1.02, 2.45] Subtotal (95% CI) 212 212 1.6% 1.46 [1.25, 1.7] Total events 153 104 Heterogeneity: $\chi^2 = 0.62$, $df = 5 (P = 0.99)$; $I^2 = 0(\%)$ Test for overall effect: $Z = 4.74$ ($P < 0.00001$) Total (95% CI) 4259 3943 100.0% 1.19 [1.17, 1.21] Total events 3831 22874 Heterogeneity: $\chi^2 = 111.71$, $df = 96$ ($P = 0.13$); $I^2 = 14\%$ Test for overall effect: $Z = 17.32$ ($P < 0.00001$)	Subtotal (95% CI)				20 3731	0.5% 98.4%	1.36 [0.97, 1.97] 1.19 [1.16, 1.21]	•
Test for overall effect: $Z = 16.86$ ($P < 0.00001$) 1.1.2 myocardial infarction induced heart failure Guo Q. 2009 22 35 17 35 0.2% 1.29 [0.85, 1.98] Mo C.R. 2002 24 36 16 38 0.2% 1.58 [1.02, 2.45] Song Q. 2001 21 23 14 22 0.3% 1.43 [1.06, 1.93] Zeng Y. 2009 22 28 12 23 0.2% 1.51 [0.97, 2.33] Zeng Y. 2009 22 28 12 23 0.2% 1.51 [0.97, 2.33] Zhang H.X. 2011 40 54 29 56 0.4% 1.43 [1.06, 1.93] Subtotal (95% CI) 212 212 1.6% 1.46 [1.25, 1.7] Total events 153 104 Heterogeneity: $\chi^2 = 0.62$, $df = 5$ ($P = 0.99$); $I^2 = 0$ (%) Test for overall effect: $Z = 4.74$ ($P < 0.00001$) Total (95% CI) 4259 3943 100.0% 1.19 [1.17, 1.21] Total events 3831 2874 Heterogeneity: $\chi^2 = 11.71$, $df = 96$ ($P = 0.13$); $I^2 = 14\%$ Test for overall effect: $Z = 1.732$ ($P < 0.00001$)	Heterogeneity: $\chi^2 = 10^4$	4.42, df =	90 (P =	0.14);	$I^2 = 1$	4(%)		
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Test for overall effect: Z	= 16.86	(P < 0.00	0001)				
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$						re 0.2%	1.29 [0.85, 1.98]	ı —
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Mo C. R. 2002 Song O. 2001	24	36	16	38	0.2%	1.58 1.02, 2.45	
Total events 153 104 Heterogeneity: $\chi^2 = 0.62, df = 5 (P = 0.99); l^2 = 0(\%)$ 104 Test for overall effect: Z = 4.74 (P < 0.00001)	Zeng Y. 2009	22	28	12	23	0.2%	1.51 [0.97, 2.33]	
Total events 153 104 Heterogeneity: $\chi^2 = 0.62, df = 5 (P = 0.99); l^2 = 0(\%)$ 104 Test for overall effect: Z = 4.74 (P < 0.00001)	Zeng Y. L. 2011 Zhang H.X. 2011		36		38	0.2%	1.45 [1.06, 1.95	
$ \begin{array}{c c} \text{Heterogeneity: } \chi^2 = 0.62, df = 5 (P = 0.99); l^2 = 0(\%) \\ \text{Test for overall effect: } Z = 4.74 (P < 0.00001) \\ \text{Total } (95\% \text{ CI}) & 4259 & 3943 100.0\% 1.19 [1.17, 1.21] \\ \text{Heterogeneity: } \chi^2 = 111.71, df = 96 (P = 0.13); l^2 = 14\% \\ \text{Heterogeneity: } \chi^2 = 17.32 (P < 0.00001) \\ \text{Test for overall effect: } Z = 17.32 (P < 0.00001) \\ \end{array} $	lotal events	153	212	104	212	1.6%	1.46 [1.25, 1.7]	-
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Heterogeneity: $\chi^2 = 0.6$	2, $df = 5$		$9); I^2 =$	0(%)			
Total events 2 3831 2874 Heterogeneity: $\lambda^2 = 111.71$, $df = 96$ ($P = 0.13$); $l^2 = 14\%$ 0.5 0.7 1 1.5 2 Test for overall effect: $Z = 17.32$ ($P < 0.00001$) 0.5 0.7 1 1.5 2		= 4.74 (1			3943	100.0%	1.19 [1.17 1.21]	
Test for overall effect: $Z = 17.32 \ (P < 0.00001)$ 0.5 0.7 1 1.5 2	Total events	3831		2874				
Test for subgroup differences: $\chi^2 = 6.67$, $df = 1$ ($P = 0.01$), $I^2 = 85\%$ Favours control Favours SFI	Test for overall effect: Z	= 17.32	(P < 0.00)	0001)				
	lest for subgroup di	tterence	s: $\chi^2 =$	6.67, df	= 1 (P = 0.01), $I^2 = 85\%$	ravours control Favours SFI

FIGURE 4: Forest plot of comparison: effect rate.

	Shenfu injection	Control	Risk ratio	Risk ratio
Study or subgroup		Events Total Weight	IV, fixed, 95% CI	IV, fixed, 95% CI
Chen Z. G. 200 Gao Z. W. 1995 Guo Y. F. 2009 Lei W. G. 2003 Liu Y. J. 2005 Luo X. Y. 2009 Wang X. M. 20 Subtotal (95% Total events Heterogeneity:	$\begin{array}{cccccccccccccccccccccccccccccccccccc$		$\begin{array}{c} 0.9 \ [0.06, 13.7] \\ 0.6 \ [0.16, 2.29] \\ 0.43 \ [0.12, 1.6] \\ 1.17 \ [0.39, 3.54] \\ 0.32 \ [0.03, 2.93] \\ 1 \ [0.06, 15.55] \\ 0.5 \ [0.05, 5.36] \\ 0.68 \ [0.36, 1.26] \end{array}$	
Guo Q. 2009 Mo C. R. 2002 Zeng Y. L. 2011 Zhang H.X. 20 Subtotal (95% Total events Heterogeneity:	8 35 9 36 8 54 11 9 36 CI) 161 - 34	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	0.67 [0.31, 1.43] 0.5 [0.26, 0.96] 0.46 [0.22, 0.97] 0.5[0.26, 0.96] 0.52 [0.37, 0.74]	* * * *
	50 $\chi^2 = 3.12, df = 10$ effect: $Z = 3.81$ (F		0.01	0.1 1 10 100 ours SFI Favours control

FIGURE 5: Forest plot of comparison: death.

Study or subgroup		u injec			ntrol SD	Tota	l Weight	Mean difference IV, random, 95% (Mean difference CI IV, random, 95% CI
Study of Subgroup	Ivicali	00	1010	Ivicali	00	1014	i weight	1, 141140111, 5570 (
Chen H. Y. 2011	525.2	168.8	30	758.6	274.3	28	7.1%	-233.4 [-351.6, -115.2]	_ - _
Chen X. B. 2009	376.5	205.9	40	766.2	297.6	40	7.3%	-389.7 [-501.85, -277.55]	
Chen Z. G. 2009	249.8	5.2	30	306.2	50	27	10.5%	-56.4 [-75.35, -37345]	=
Guo J.J. 2006	644.5	285.4	36	848.5	352.3	26	5.4%	-204 [-368.41, -39.59]	
Guo Q. 2009	186.8	41.9	35	291.7	55.6	35	10.5%	-104.9 [-127.96, -81.84]	T
Liu S. M. 2008	382.1	151.8	74	467.4	171.2	78	9.7%	-85.3 [-136.68, -33.92]	
Luo X. Y. 2009	1.796.5	237.7	50	2.296.5	263.5	50	7.9%	-500 [-598.36, -401.64]	
Lv G. 2010	251.4	63.3	31	424.3	47.5	30	10.4%	-172.9 [-200.93, -144.87]	+
Tan L. J. 2011	212.5	56.7	38	357.6	73.4	30	10.3%	-145.1 [-176.96, -113.24]	.
Wu. H. J. 2009	512.9	176	33	553.9	150.4	29	8.6%	-41 [-502, -37345]	
Yang Z. Y. 2008	375.6	204.2	40	765.9	297	40	7.3%	-390.3 [-502, -278.6]	_ _
Zhou G. 2010	576.5	201.4	19	887.4	322.6	18	5.1%	-310.9 [-485,29, -136.51]	
Total (95% CI)			456			431	100.0%	-201.26 [-255.27, -147.25]	▲ · · · ·
									-500-250 0 250 500
2		2							

Heterogeneity: $\tau^2 = 7116.18$; $\chi^2 = 176.82$; df = 11 (P < 0.00001); $I^2 = 94\%$ Test for overall effect: Z = 7.3 (P < 0.00001)

Favours SFI Favours control

FIGURE 6: Forest plot of comparison: NT-proBNP.

western medicine may sometimes fail to treat an illness, whereas such illness is reportedly improved by the so-called complementary medicine based on a different theory [110, 111]. Although conventional therapeutic approaches were used in HF, it remained a cardiovascular disease with an increasing hospitalization burden and an ongoing drain on health care expenditures [2]. TCM plays an important role in treating HF in China. SFI was a traditional Chinese Patent Medicine based on TCM theory, which was approved by the Chinese State Food and Drug Administration. In recent 10 years, it has been widely used for HF in many hospitals and clinics. However, few RCTs of SFI were reported in English journals, and it was difficult for western doctors to accept SFI as an alternative medicine. Although there were two systematic reviews about SFI for HR published in Chinese journal [112, 113], only 16 and 8 trials were included in their study. Therefore, the present study aimed to systematically assess the efficacy and safety of SFI for HR.

Data from the 97 RCTs demonstrated that SFI combined with conventional medication may be more effective on HF than conventional medication only. With improvement of cardiofunction of patients, based on NYHA Classification of

	Shenf	u injec	ction	Сс	ontro	ol		Mean difference	Mean difference
Study or subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, fixed, 95% CI	IV, fixed, 95% CI
Chen X. B. 2009	443	66	40	395	64	40	1.9%	48 [19.51, 76.49]	
Chen X. L. 2009	250	112	35	200	90	35	0.7%	50 [2.4, 97.6]	
Guo J.J. 2006	497	74	36	413	67	26	1.2%	84 [48.68, 119.32]	
Huang T. 2009	329	64	38	280	52	38	2.2%	49 [22.78, 75.22]	
Liu S. S. 2007	218	17	50	211	15	50	38.6%	7 [0.72, 13.28]	-
Ma H. W. 2005	216	18	42	203	16	42	28.7%	13 [5.72, 20.28]	+
Yang Z. Y. 2008	445	65	40	395	63	40	1.9%	50 [21.95, 78.05]	— ,
Zhan L. S. 2001	330	18	46	316	17	32	24.7%	14 [6.14, 21.86]	*
Total (95% CI)			327			303	100%	14.22 [10.31, 18.13]	
Heterogeneity: $\chi^2 =$					2 =	83%			-100 -50 0 50 100
Test for overall effect	Z = 7.1	4 (P <	0.0000	1)					Favours control Favours SFI

FIGURE 7: Forest plot of comparison: 6-MWD.

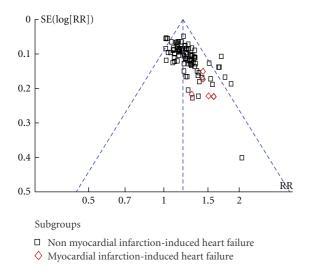


FIGURE 8: Funnel plot of comparison: effect rate.

Clinical Status and Killip's Rating Standards, the effect rate of SFI group was, on average, 17 percent more than control group (RR, 1.19; 95% CI, 1.17 to 1.21). Mortality data was another primary outcome. In eleven trials in which death was recorded, meta-analysis showed that mortality was significantly lower in SFI group than control group. This result was mainly contributed by subgroup of HF induced by myocardial infarction, for patients in this subgroup were more vulnerable.

Ultrasonic cardiography is widely used in inspection for HF patients. From results of ultrasonic cardiography, the systolic and diastolic functions of heart can be interpreted. LVEF, CO, CI, SV, LVDd, and E/A were reviewed by us, respectively. There was significant difference between SFI group and control group in all of the outcomes except LVDd. Since SV, CO, CI, and LVEF indicate heart systolic function, and E/A indicate heart diastolic function, conclusion can be drawn that SFI benefits both systolic and diastolic functions of heart. But it did not have significant effect on expansion of heart. NT-proBNP level in serum of SFI group was significantly lower than the control group, which is inconsistent with effect rate. 6-MWD results of patients of SFI group also are better than thos of control group. It indicates that SFI had a tendency to improve life status. Furthermore, heart rate was obviously reduced in SFI group, which could be related to alleviation of HF.

Meta-analysis on LVEF, CO, CI, SV, LVDd, E/A, heart rate, and NT-proBNP all showed significant heterogeneity. Several possible explanations can be given, for example, different complications, different instruments employed for test, and difference in methodological rigor.

However, we should consider the following limitations before accepting the findings of this paper.

Firstly, the methodological quality of the included studies is generally poor. Although all trials claimed to perform randomization, only eleven trials reported the procedure to generate the sequence, while the rest of trials did not give any details of the randomization method. Thus, whether randomization was effectively conducted in these trials was doubtful. Blinding was mentioned in four trials, with one trial blinded patients and outcome assessors [43] and three blinded patients only [44-46]. Neither of them described the methods of allocation concealment. Dropouts account and intention to treat analysis were not mentioned in all the trails. Due to inadequate reporting of methodological design, it was possible that there was performance bias and detection bias due to patients and researchers being aware of the therapeutic interventions for the subjective outcome measures. Therefore, we cannot draw a confident conclusion that there were significant beneficial effects of SFI combined with conventional medicine treatment compared with conventional medicine treatment.

Secondly, limited outcomes were reported, especially death and adverse events. Since HF is a disease with high mortality, death is the most important primary outcome. However, only eleven studies out of ninety seven trials reported death, and most of the eleven trials assessed mortality at the end of treatment, without followup. Another outcome was adverse events, to which more attention should be attached. Only 37.4% of the trials described the occurrence of adverse events, indicating an incomplete evaluation of the safety profile of SFI, as well as poor quality of reporting. In most trials, the duration of therapy and followup was

too short to achieve conclusive results, except that only one trial had a treatment of 10 months [47]. Only 6 included trials had a followup period (ranged from 3 to 12 months), while in rest of studies, the outcomes were evaluated at the end of the treatment (mostly range from 14 to 21 days). In order to evaluate drug efficacy for chronic HF, long-term improvement (at least 6 months) of chronic HF-specific clinical symptoms is needed [114], because some drugs have shown to increase mortality in the long-term application despite a short-term improvement in clinical symptoms [115]. In addition, long-term toxicity assessment was also important for drug safety evaluation.

Next, although irrespective of languages, all the trials included in this paper were published in Chinese journals, Zhang et al. and Liu et al. [115, 116] found that some Asian countries including China unusually publish high proportions of positive results. Wu et al. [117] and Jin et al. [118] accounted that RCTs in Chinese journals often had problems of low methodological quality and selective publication of positive results. Considering that all of the ninety seven trials were published in Chinese, the publication bias possibly existed.

Additionally, none of the ninety seven trials reported sample size calculation, and in most trials, the sample size was limited. Further high-quality studies with larger sample size are needed to confirm the effectiveness of SFI in treating HF. Quality of life was not reported in all the including trials. Although 6-MWD showed a tendency of SFI to improve life status for HF patients, we advise future RCTs to select outcomes of life quality according to international practice.

Considering that there was no sufficient amount of highquality trials on SFI treating patients with HF, the effectiveness and safety of SFI need further rigorous trials to prove, which should be consistent with the CONSORT statement on the reporting of the results of randomized trials (http:// www.consort-statement.org/).

5. Conclusion

The preliminary conclusion of the current study suggests that SFI might be beneficial to patients with HF. More rigorously designed trails with high methodological quality are necessary for further proof.

Conflict of Interests

The authors declare that there is no conflict of interests.

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