Review Article

The Effectiveness and Safety of Huangqi Xixin Decoction for Cough Variant Asthma: A Systematic Review and Meta-Analysis

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Objective. A comprehensive and systematic review is needed to evaluate the safety and effectiveness of Huangqi Xixin decoction (HQXXD) for cough variant asthma (CVA). In this systematic review, we comprehensively interrogate the safety and effectiveness of HQXXD for CVA. Methods. An overall search for studies in main English and Chinese electronic databases from their inception to June 30, 2022, was performed. Randomized controlled trials (RCTs) involving HQXXD for CVA were included. According to Cochrane Reviewer’s Handbook, the risk of bias related to the included studies was evaluated. A meta-analysis using RevMan 5.4 software from the Cochrane Collaboration was used to integrate the outcomes of the included RCTs. Results. A systematic review and meta-analysis were conducted using the seven eligible RCTs that had been retrieved. The included RCT-related risk of bias was evaluated. According to the findings of the meta-analysis, the HQXXD group had significantly higher total effective rates of clinical efficacy and airway responsiveness, and a significantly lower recurrence rate in comparison with the conventional Western medicine treatment group. Conclusion. In the treatment of CVA patients, HQXXD is safe and effective, which benefits clinical efficacy and airway responsiveness, reduces the recurrence rate, and has no adverse effects.

1. Introduction

Cough variant asthma (CVA), the primary or only clinical manifestation of chronic cough, is a particular type of asthma [1]. As with asthma, the pathogenesis of CVA is not fully clear, and its pathophysiological changes are characterized by airway hyperresponsiveness and chronic inflammation [2]. CVA is a key pathogenic pathway for chronic cough and has been the most prevalent cause in China and Japan and the second in Korea, respectively [3–5]. Currently, the guidelines have suggested that regular inhaled corticosteroids and long-acting β2-agonists can be used as the recommended treatment modalities for patients with CVA [6, 7]. However, the long-term treatment effects of some CVA patients are not ideal, the recurrence rate is still high, and adverse reactions such as allergic reactions, as well as mental or neurological reactions, may occur [8, 9]. The main syndrome types of CVA in traditional Chinese medicine (TCM) theory are characterized by Qi deficiency and vigorous wind, which are generally classified as wind cough, stubborn cough, or wheezing cough [10]. Nowadays, combined with the TCM theory, TCM prescription has achieved a certain curative effect in the treatment of CVA [11, 12].

Huangqi Xixin decoction (HQXXD) is modified from the TCM prescriptions, Zhi Soupowders, and Yu Ping Feng powders [13]. It primarily comprises Huang Qi, Xi Xin, Jing Jie, Fang Feng, Huang Qin, Bai Zhu, Fu Ling, Chan Tui, Ban Xia, Bai Bu, and Gan Cao, whose pharmaceutical Latin names are Radix Astragali (RA), Herba cum Radix Asari...
(HRA), Herba Schizonepetae (HS), Radix Saposhnikoviae (RS), Radix Scutellariae, Rhizoma Atractylodis macrocephalae, Poria, Periostracum cicadae, Rhizoma Pinelliae, Radix Stemonae, and Radix Glycyrrhizae, respectively. RA, HRA, HS, and RS are the main active herbs in HQXXD, which are considered monarch and minister herbs according to the TCM theory. RA could tonify lung Qi; HRA, HS, and RS could tonify the spleen and stomach and dispel wind dampness to strengthen the effect of RA [14].

An earlier systematic review reported that there was no significant difference between the control and HQXXD groups in terms of the clinical efficacy reported by the total effective rate [13]. Our latest study, which included a meta-analysis of HQXXD’s clinical curative effect on CVA, revealed that HQXXD acted on CVA in a variety of mechanisms, including through several compounds, targets, and pathways [14]. To date, numerous clinical studies using HQXXD for CVA have been published [15, 16]. However, the indicators evaluated were not comprehensive, such as recurrence rate, pulmonary function indices, and biochemical test indices. Therefore, a comprehensive and systematic review is needed to analyze the safety and effectiveness of HQXXD for CVA. This study aimed to comprehensively assess the safety and effectiveness of HQXXD for CVA.

2. Materials and Methods

2.1. Protocol and Registration. The protocol of the review was registered on the PROSPERO platform (https://www.crd.york.ac.uk/PROSPERO/) with the registration number CRD42021235772. PROSPERO, produced by the Centre for Reviews and Dissemination (CRD) and funded by the National Institute for Health Research (NIHR), is a global database of prospectively registered systematic reviews. At the beginning of the registration, PROSPERO can compare the completed review with the content of the protocol by offering a comprehensive listing of systematic reviews, to avoid the circumstances of duplication and reporting bias [17, 18]. Meanwhile, the review protocol has been published in an open-access journal [19].

The Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) 2020 statement served as the basis for our review [20, 21]. The PRISMA 2020 checklist is shown in Supplementary File 1.

2.2. Ethical Consideration. This systematic review collected data from open databases. All the eligible studies were approved by the local institutional ethics committee, and the written informed consent of the participants was collected. This systematic review did not directly involve the patient’s privacy, so additional ethical approval was not necessary.

2.3. Search Strategy. We performed an overall search for published studies in main English and Chinese electronic databases from their inception to June 30, 2022, which include MEDLINE via PubMed, EMBASE via Ovid, Cochrane Central Register of Controlled Trials (CENTRAL), Chinese National Knowledge Infrastructure (CNKI), Chongqing VIP information (CQVIP), Wanfang database, and Chinese Biomedical Database (CBM). We tried to contact the authors to obtain the data we needed. The references in the included literature and systematic reviews were also inspected. On the basis of the databases feature, we modulated the search strategies of title, abstract, or keywords. Search strategies in PubMed are presented in Supplementary File 2, and the keywords “cough variant asthma” combined with “Huangqi Xixin decoction”; “Radix Astragali”; “Herba cum Radix Asari”; “Herba Schizonepetae”; “Radix Saposhnikoviae”; “traditional Chinese Medicine”; “Chinese Medicine”; and “herbal medicine” were used for the search. Some keywords were adjusted slightly in searching on different electronic databases.

2.4. Eligibility Criteria. The eligibility criteria strictly complied with PICOS (participant, intervention, comparison, outcome, and study design) principles.

2.4.1. Study Design. Randomized controlled trials (RCTs) examining the safety and effectiveness of HQXXD for CVA were included in this review. We excluded case reports, retrospective observational studies, and animal research.

2.4.2. Participants. Regardless of gender, age, race, educational level, and economic and marital status, individuals diagnosed with CVA using clearly defined or internationally recognized criteria were included.

Individuals with other respiratory conditions (bronchiectasis, chronic obstructive pulmonary disease), or severe liver, kidney, or heart disease were excluded.

2.4.3. Intervention. Patients treated with HQXXD were included. HQXXD prescription can be modified, but it must contain monarch and minister herbs (RA, HRA, HS, and RS). Eligible treatments could be employed as monotherapy or combined conventional Western medicine treatment (CWMT).

2.4.4. Comparison. Comparators included placebo, CWMT, or no interventions. Studies comparing other TCM prescription treatments in comparison were excluded.

2.4.5. Outcomes. All the outcome indicators from the included studies were retrieved and evaluated. We performed a meta-analysis when the indicators could be subjected to a meta-analysis and a descriptive analysis when meta-analysis could not be done. The main outcomes included clinical efficacy, airway reactivity, recurrence rate, pulmonary function indices, biochemical test indices, and adverse events.
2.5. Study Selection, Data Extraction, and Quality Assessment

2.5.1. Study Selection. All the literature was retrieved independently by three reviewers (Wang, Xia, and Hu). The initially identified references from searching databases were exported to NoteExpress 3.2.0 software. The duplicate studies were eliminated using NoteExpress software duplication models [22]. After that, the three reviewers independently evaluated each study’s titles, abstracts, and keywords to screen possible eligible studies according to the predefined evaluation criteria. Finally, the three reviewers scrutinized and cross-checked the full text of previous eligible studies in the second stage and confirmed the final included studies for this systematic review. The reasons for excluding each study were recorded in detail in the second stage. A discussion was done to resolve the disagreements of the three reviewers at any stage of the selection process. Another reviewer (Zhang) was invited as an arbitrator to make a final decision when the disagreements could not be resolved after the discussion among the three reviewers. A PRISMA-compliant flow chart was used to outline the identification and selection procedures for this systematic review.

2.5.2. Data Extraction and Dealing with Missing Data. Three reviewers (Wang, Xia, and Hu) conducted the data extraction procedure independently after completing a standard data extraction sheet. They cross-checked these results and examined whether there were any differences. A discussion was done to resolve the inconsistent opinions of the three reviewers. Another reviewer (Zhang) was invited as an arbitrator to judge these disagreements. The following list of information extracted from the original articles was saved in a standard data extraction sheet: (1) title, study year, country, and authors; (2) study design; (3) methodology: randomization, allocation concealment, patient and assessor blinding, inadequate outcome data, selective outcome reporting, and other risks of bias; (4) sample size; (5) age and gender in each group; (6) diagnostic criteria; (7) intervention group (methods of treatment, compositions of HQXXD prescription, duration of treatment); (8) comparison group (methods of treatment, duration of treatment); (9) main outcomes; and (11) adverse events. We contacted the corresponding authors of each study through email when the above information was missing.

2.5.3. Quality Assessment. The quality of each eligible study was independently evaluated by three reviewers (Wang, Xia, and Hu) employing the Cochrane risk-of-bias tool from the Cochrane Handbook V.5.1.0. In addition to patient and assessor blinding, random sequence generation, allocation concealment, inadequate outcome data, selective outcome reporting, and other bias risks were among the seven criteria. Each item was categorized as having a low, high, or unclear risk of bias [23, 24]. In terms of other biases, we also carefully assessed the baseline imbalance and different sources of funding support. Disagreements were discussed with another reviewer (Zhang).

2.6. Statistical Analysis and Assessment of Heterogeneity. We used Review Manager (RevMan) 5.4 software to perform statistical analysis. To conduct a meta-analysis, we calculated the odds ratio (OR) with a 95% confidence interval (CI) for dichotomous data and the mean difference (MD) with a 95% CI for continuous data. The chi-square and I^2 tests were performed to explore the heterogeneity. When I^2 < 50% and P > 0.1, there was homogeneity between each study. For the meta-analysis, the Mantel–Haenszel (M-H) fixed-effects model was employed. Otherwise, when I^2 ≥ 50% or P < 0.1, the included studies were considered heterogeneous. Initially, to explore possible factors affecting the clinical heterogeneity, we re-assessed the demographic characteristics of the CVA patients and the variation of interventions between each included study. When clinical heterogeneity existed in this systematic review, we performed descriptive analysis. Otherwise, a random-effects model based on an inverse variance statistical approach was employed for further meta-analysis after getting rid of the clinical heterogeneity [25, 26]. When we performed the meta-analysis, P < 0.05 was taken as a significant value.

2.7. Subgroup Analysis. When heterogeneity was high and the number of included studies was sufficient, we assessed different interventions (monotherapy or combined CWMT), different control (placebo, CWMT, or no interventions), and different durations of treatment to carry out subgroup analysis [27].

2.8. Sensitivity Analysis. Based on methodological quality (low-quality studies were omitted to re-evaluate the results of this meta-analysis), statistical model (random-effects or fixed-effects model was used for analysis), and sample size (studies with smaller sample size were omitted to re-evaluate the outcomes of this meta-analysis), sensitivity analysis was carried out [28].

2.9. Publication Bias. When there were over five studies included in the meta-analysis, we used RevMan 5.4 software to assess publication bias with a funnel plot [29].

2.10. Summary of Evidence. We used the GRADEpro Web tool (https://gradepro.org/) to evaluate the quality of each main outcome in this systematic review and categorized them into 4 grades: high, medium, low, or very low. GRADEpro conducts the guideline development progression sternly depending on the GRADE methodology, including multiple fields of evidence such as summarization, recommendations, and dissemination. The judgments were based on the risk of bias, inconsistency, imprecision,
indirectness, large effect, dose-response gradient, publication bias, and plausible confounding [30].

3. Results

3.1. Study Selection. After excluding 3727 studies from the retrieved 5687 studies via database searching, because of the faint relevance expressed from the title and the abstract, 1932 were also removed from the remnant studies, and therefore, 28 studies were retained. Following the exclusion process of 21 studies due to various reasons, the systematic review eventually included seven RCTs [15, 16, 31–35]. In Figure 1, the findings of the literature screening and the process are displayed.

3.2. Description of Included Studies. Seven eligible RCTs were screened. All seven RCTs involving 512 patients were carried out in China. All of them were single-center studies. Participants in one study [16] were children, and participants in the other studies were adults. All studies used the prescription of HQXXD as monotherapy. The control group included CWMT; one study [15] used terbutaline sulfate tablets; one study [34] used procaterol hydrochloride tablets; three studies [31–33] used procaterol hydrochloride tablets and salbutamol aerosol; one study [35] used ambroxol hydrochloride tablets and terbutaline sulfate tablets; and one study [16] used montelukast sodium chewable tablets and budesonide aerosol. Table 1 lists the basic features of the studies that were included, and Table 2 lists the compositions of HQXXD prescriptions used in each study’s experimental group.

3.3. Methodological Quality. Two RCTs utilized adequate methods of random sequence generation, one [32] introduced the envelope method and another [35] introduced the draw random method, and other studies did not specifically describe the randomized methods and allocation concealment; none RCT introduced blindness; two RCT [31, 34] had

Figure 1: Flow diagram for the research selection process. Through database search, 5687 studies in total were retrieved. Finally, the systematic evaluation involved seven RCTs.
incomplete outcome data; and selective reports cannot be identified from all studies. The risk-of-bias graph about each risk-of-bias item presented as percentages across all included studies is shown in Figure 2. In addition, the risk-of-bias summary about each risk-of-bias item for each included study is shown in Figure 3 and Table S1.

### 3.4. Outcomes

Six RCTs [15, 16, 32–35] compared the clinical efficacy reported by total effective rate; three RCTs [31, 32, 35] compared the total effective rate of airway responsiveness, and three RCTs [32–34] compared the recurrence rate, but one RCT [34] had incomplete data; two RCTs [31, 32] compared the improvement rate of cough, throat itching, and cough up phlegm, but one RCT [31] had incomplete data; one RCT [16] compared the serum tumor necrosis factor α (TNF-α), interleukin-8 (IL-8), and IL-6 indices, and pulmonary function indices including forced expiratory flow at 50% of forced vital capacity (FEF50), forced expiratory flow at 75% of forced vital capacity (FEF75), and maximal mid-expiratory flow (MMEF75/25). Adverse reactions were involved in the five studies [15, 32–35] with no adverse reactions reported, whereas the other two studies [16, 31] did not probe into adverse reactions. Table 3 provides a summary of the key results.

| Study year [ref] | Country | Sample size (experimental/control) | Mean age (years) (experimental/control) | Experimental | Control | Duration |
|------------------|---------|-----------------------------------|----------------------------------------|--------------|---------|----------|
| Wang and Xie 2009 [15] | China | 42 (21/21) | 34.5 ± 2.7/33.2 ± 2.6 | HQXXD | Terbutaline sulfate tablets | 3 weeks |
| Zhao 2014 [31] | China | 90 (45/45) | 42.1 ± 8.27/39.1 ± 11.30 | HQXXD | Procaterol hydrochloride tablets + salbutamol aerosol | 4 weeks |
| Fan and Xie 2014 [32] | China | 60 (30/30) | 42.10 ± 8.27/38.2 ± 10.35 | HQXXD | Procaterol hydrochloride tablets + salbutamol aerosol | 4 weeks |
| Li 2015 [33] | China | 70 (35/35) | 50 ± 1.5/49 ± 1.3 | HQXXD | Procaterol hydrochloride tablets + salbutamol aerosol | 4 weeks |
| Wang and Xie 2015 [34] | China | 90 (45/45) | 41.12 ± 7.24/40.18 ± 9.35 | HQXXD | Procaterol hydrochloride tablets | 4 weeks |
| Wei and Qin 2019 [35] | China | 80 (40/40) | 45.35 ± 3.88/43.25 ± 3.78 | HQXXD | Ambroxol hydrochloride tablets + terbutaline sulfate tablets | 4 weeks |
| Wang 2020 [16] | China | 80 (40/40) | 4.56 ± 2.28/4.62 ± 2.13 | HQXXD | Montelukast sodium chewable tablets + budesonide aerosol | 8 weeks |

RCT: randomized controlled trial, HQXXD: Huangqi Xixin decoction, and CVA: cough variant asthma.

### 3.5. Meta-Analysis

#### 3.5.1. Subgroup Analysis and Sensitivity Analysis

We first assessed the heterogeneity based on interventions, controls, and duration of treatment. When the heterogeneity was low, then subgroup analysis was not carried out. Sensitivity analysis was performed based on methodological quality, statistical model, and sample size. The sensitivity analysis demonstrated that the robustness and reliability of the pooled results were fair.

#### 3.5.2. Total Effective Rate of Clinical Efficacy

A total of 422 patients were included in the six studies [15, 16, 32–35] that compared the total effective rate of clinical efficacy with 211 in the experimental group and 211 in the control group. The heterozygosity test revealed that there was no heterogeneity in the six studies ($P = 0.77, I^2 = 0\%$). When OR values were combined using the fixed-effects model, the pooled OR was 3.45 (95% CI [1.78–6.67], $P < 0.0002$). These findings demonstrated that the total effective rate of clinical efficacy in the experimental group was substantially higher than that of the control group (Figure 4).

#### 3.5.3. Total Effective Rate of Airway Responsiveness

There were 200 individuals in total in the three studies [31, 32, 35] that compared the total effective rate of airway responsiveness, with 100 in the experimental group and 100 in the control group, respectively. The heterozygosity test revealed that there was no heterogeneity between the two studies ($P = 0.87, I^2 = 0\%$). When OR values were combined using the fixed-effects model, the pooled OR was 5.11 (95% CI [1.83–14.25], $P < 0.002$). This showed that the total effective rate of airway responsiveness in the experimental group was considerably higher than that in the control group (Figure 5).

#### 3.5.4. Recurrence Rate

A total of 130 patients were included in the two studies [3, 33] that compared the recurrence rate, 65 of whom were in the experimental group and 65 in the control group, respectively. The heterozygosity test revealed that there was no heterogeneity between the two studies ($P = 0.53, I^2 = 0\%$). The pooled OR was 0.20 (95% CI [0.09–0.44], $P < 0.0001$) when the fixed-effects model was applied to combine OR values. This showed that the recurrence rate in the experimental group was considerably lower than that in the control group (Figure 6).
### Table 2: Compositions of TCM prescriptions.

| Study year (ref) | TCM prescriptions | Compositions of TCM prescriptions |
|------------------|-------------------|----------------------------------|
| Wang and Xie 2009 [15] | HQXXD | Radix *Astragali* (Latin name: *Astragalus* root) | Huang Qi (Chinese name: 黄芪) |
|                   |                   | Herba cum Radix *Asari* (Latin name: *Asarum* Xi Xin) | Xi Xin (Chinese name: 朮) |
|                   |                   | Radix Schizonepetae (Latin name: Schizonepeta stem or bud) | Jing Jie (Chinese name: 茜草) |
|                   |                   | Radix Saposhnikoviae (Latin name: Saposhnikoviae root) | Fang Feng (Chinese name: 茜草) |
|                   |                   | Ramulus *Cinnamomi* (Latin name: Cinnamon twig) | Gui Zhi (Chinese name: 肉桂) |
|                   |                   | Rhizoma *Atractylodis macrocephalae* (Latin name: Atractylodis rhizome) | Bai Zhu (Chinese name: 苍术) |
|                   |                   | Periostracum cicaeae (Latin name: Cicada molting (slough)) | Chan Tui (Chinese name: 蝉蜕) |
|                   |                   | Rhizoma Pinelliae (Latin name: Pinellia rhizome) | Ban Xia (Chinese name: 丹参) |
|                   |                   | Radix Stemonae (Latin name: Stemona root) | Bai Bu (Chinese name: 丹参) |
|                   |                   | Radix Glycyrrhizae (Latin name: Licorice root) | Gan Cao (Chinese name: 甘草) |
| Zhao 2014 [31]    | HQXXD | Radix *Astragali* (Latin name: *Astragalus* root) | Huang Qi (Chinese name: 黄芪) |
|                   |                   | Herba cum Radix *Asari* (Latin name: *Asarum* Xi Xin) | Xi Xin (Chinese name: 朮) |
|                   |                   | Radix Schizonepetae (Latin name: Schizonepeta stem or bud) | Jing Jie (Chinese name: 茜草) |
|                   |                   | Radix Saposhnikoviae (Latin name: Saposhnikoviae root) | Fang Feng (Chinese name: 茜草) |
|                   |                   | Radix Scutellariae (Latin name: Scute) | Huang Qin (Chinese name: 荆芥) |
|                   |                   | Rhizoma *Atractylodis macrocephalae* (Latin name: Atractylodis rhizome) | Bai Zhu (Chinese name: 苍术) |
|                   |                   | Periostracum cicaeae (Latin name: Cicada molting (slough)) | Chan Tui (Chinese name: 蝉蜕) |
|                   |                   | Rhizoma Pinelliae (Latin name: Pinellia rhizome) | Ban Xia (Chinese name: 丹参) |
|                   |                   | Radix Stemonae (Latin name: Stemona root) | Bai Bu (Chinese name: 丹参) |
|                   |                   | Radix Glycyrrhizae (Latin name: Licorice root) | Gan Cao (Chinese name: 甘草) |
| Fan and Xie 2014 [32] | HQXXD | Radix *Astragali* (Latin name: *Astragalus* root) | Huang Qi (Chinese name: 黄芪) |
|                   |                   | Herba cum Radix *Asari* (Latin name: *Asarum* Xi Xin) | Xi Xin (Chinese name: 朮) |
|                   |                   | Radix Schizonepetae (Latin name: Schizonepeta stem or bud) | Jing Jie (Chinese name: 茜草) |
|                   |                   | Radix Saposhnikoviae (Latin name: Saposhnikoviae root) | Fang Feng (Chinese name: 茜草) |
|                   |                   | Radix Scutellariae (Latin name: Scute) | Huang Qin (Chinese name: 荆芥) |
|                   |                   | Rhizoma *Atractylodis macrocephalae* (Latin name: Atractylodis rhizome) | Bai Zhu (Chinese name: 苍术) |
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|                   |                   | Radix Stemonae (Latin name: Stemona root) | Bai Bu (Chinese name: 丹参) |
|                   |                   | Radix Glycyrrhizae (Latin name: Licorice root) | Gan Cao (Chinese name: 甘草) |
| Li 2015 [33]      | HQXXD | Radix *Astragali* (Latin name: *Astragalus* root) | Huang Qi (Chinese name: 黄芪) |
|                   |                   | Herba cum Radix *Asari* (Latin name: *Asarum* Xi Xin) | Xi Xin (Chinese name: 朮) |
|                   |                   | Radix Schizonepetae (Latin name: Schizonepeta stem or bud) | Jing Jie (Chinese name: 茜草) |
|                   |                   | Radix Saposhnikoviae (Latin name: Saposhnikoviae root) | Fang Feng (Chinese name: 茜草) |
|                   |                   | Radix Scutellariae (Latin name: Scute) | Huang Qin (Chinese name: 荆芥) |
|                   |                   | Rhizoma *Atractylodis macrocephalae* (Latin name: Atractylodis rhizome) | Bai Zhu (Chinese name: 苍术) |
|                   |                   | Periostracum cicaeae (Latin name: Cicada molting (slough)) | Chan Tui (Chinese name: 蝉蜕) |
|                   |                   | Rhizoma Pinelliae (Latin name: Pinellia rhizome) | Ban Xia (Chinese name: 丹参) |
|                   |                   | Radix Stemonae (Latin name: Stemona root) | Bai Bu (Chinese name: 丹参) |
|                   |                   | Radix Glycyrrhizae (Latin name: Licorice root) | Gan Cao (Chinese name: 甘草) |
| Wang and Xie 2015 [34] | HQXXD | Radix *Astragali* (Latin name: *Astragalus* root) | Huang Qi (Chinese name: 黄芪) |
|                   |                   | Herba cum Radix *Asari* (Latin name: *Asarum* Xi Xin) | Xi Xin (Chinese name: 朮) |
|                   |                   | Radix Schizonepetae (Latin name: Schizonepeta stem or bud) | Jing Jie (Chinese name: 茜草) |
|                   |                   | Radix Saposhnikoviae (Latin name: Saposhnikoviae root) | Fang Feng (Chinese name: 茜草) |
|                   |                   | Radix Scutellariae (Latin name: Scute) | Huang Qin (Chinese name: 荆芥) |
|                   |                   | Rhizoma *Atractylodis macrocephalae* (Latin name: Atractylodis rhizome) | Bai Zhu (Chinese name: 苍术) |
|                   |                   | Periostracum cicaeae (Latin name: Cicada molting (slough)) | Chan Tui (Chinese name: 蝉蜕) |
|                   |                   | Rhizoma Pinelliae (Latin name: Pinellia rhizome) | Ban Xia (Chinese name: 丹参) |
|                   |                   | Radix Stemonae (Latin name: Stemona root) | Bai Bu (Chinese name: 丹参) |

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3.5.5. Publication Bias Analysis. Utilizing funnel plots, the publication bias was investigated. We created funnel plots showing the clinical efficacy as presented as the total effective rate, where the horizontal coordinate was each outcome OR value and the longitudinal coordinate was the SE (log [OR]). The funnel plots are shown in Figure 7.

3.6. Safety of HQXXD for CVA. Five studies [15, 32–35] involved adverse reactions and reported that HQXXD had no adverse reaction in treatment. Other studies [16, 31] did not probe into the adverse reactions. HQXXD is safe for CVA patients.

3.7. GRADE Level of Evidence. The GRADE levels of evidence were moderate for total effective rates of clinical efficacy and airway responsiveness outcomes. For the outcome of recurrence rate, the GRADE level of evidence was low. The GRADE evidence profiles are shown in Figure 8. The high risk of bias was the primary cause of a decreasing level.

4. Discussion

Based on tonifying lung Qi and dispelling wind dampness, TCM has been used in the treatment of CVA and the treatment effect is clear [12, 36]. HQXXD, in which RA, RS, HRA, and HS are monarch and minister herbs, could tonify lung Qi and dispel wind dampness to treat CVA according to the TCM theory [13]. On the basis of a network pharmacology approach, a previous study interrogated the multicomponent, multtarget, and multi-pathway characteristics of HQXXD acting on CVA [14]. Basic and clinical study of HQXXD treating CVA deserves further research.

However, there is no systematic review showing comprehensive evidence regarding the safety and effectiveness of HQXXD in CVA. This systematic review illustrated that total effective rates of clinical efficacy and airway responsiveness were considerably higher in the HQXXD group in comparison with those in the CWMT group (both had $P < 0.05$); the recurrence rate was considerably lower in the HQXXD group in comparison with that in CWMT group ($P < 0.05$).

The total effective rate of clinical efficacy is the most intuitive index of clinical efficacy evaluation. When comparing the total effective rate of clinical efficacy between the HQXXD and the CWMT groups, individual studies [15, 32–35] did not find a considerable difference. In the previous meta-analyses, one study [13] showed no significant difference, whereas one study [14] showed a statistically significant difference. Based on expanding the sample size, this systematic review confirmed that the total effective rate of clinical efficacy in the HQXXD group was substantially higher than that of the CWMT group.

Airway responsiveness is an important pathophysiological mechanism in CVA [37]. When comparing the total effective rate of airway responsiveness between the HQXXD and the CWMT groups, individual studies [31, 32, 35] did not find a statistically significant difference. This systematic review clarified that the HQXXD group had a substantially
higher total effective rate of airway responsiveness than the CWMT group.

Through CWMT, the long-term treatment effects of some CVA patients are not ideal and the recurrence rate is still high [8, 9]. There is no systematic review regarding the recurrence rate of HQXXD for CVA patients. This systematic review was the first to report that the recurrence rate in the HQXXD group was substantially lower than that in the CWMT group.

Regarding the safety of HQXXD for CVA, five studies [15, 32–35] regarding adverse reactions showed that HQXXD had no adverse reaction in treatment. Other studies [16, 31] did not involve adverse reactions. We did a descriptive analysis that HQXXD is safe for CVA patients.

Moreover, two RCTs [31, 32] compared the improvement rate of cough, throat itching, and cough up phlegm; one RCT [16] compared serum TNF-α, IL-8, and IL-6 indices, and FEF50, FEF75, and MMEF75/25. These provide references for our in-depth research and need more relevant studies to conduct a further evaluation.

This systematic review is aimed at highlighting the HQXXD prescription for routine CNA treatment, or providing a basis for further real-world RCT research of HQXXD and obtaining more evidence-based medical evidence for better application of TCM. Our review protocol was registered on the PROSPERO platform, which is produced by CRD and funded by the NIHR, aiming to avoid the circumstances of duplication and reporting bias. On the other hand, this review was performed based on the PRISMA, and all of these make our review more standardized and reliable.

One of the seven studies [16] selected for final analysis includes only children. Our inclusion and exclusion criteria did not limit the age, so it was eventually included. We performed heterogeneity analysis, which showed no obvious heterogeneity, and the number of included studies was less, so we did not conduct subgroup analysis. With the publication of more relevant RCT, further subgroup analysis may be possible to obtain more accurate evidence.

However, this systematic review may have some potential limitations. First of all, HQXXD prescription originates in China, and the results of this review may limit to Asian patients although the internationalization of TCM is becoming increasingly extensive. Secondly, some of the studies do not introduce allocation concealment and contain unclear random methods; the studies lack the introduction
### Table 3: Main outcomes of included RCTs.

| Study year (ref)          | Main outcomes                          | Main results (effect size) | Adverse events |
|--------------------------|----------------------------------------|----------------------------|----------------|
| Wang and Xie 2009 [15]   | Total effective rate of clinical efficacy | OR 6.25 [0.66, 59.03]     | No adverse reaction |
| Zhao 2014 [31]           | Improvement rate of cough               |                            |                |
|                          | Improvement rate of throat itching      |                            |                |
|                          | Improvement rate of cough up phlegm     |                            |                |
|                          | Total effective rate of airway responsiveness | OR 4.26 [0.81, 22.53]   |                |
| Fan and Xie 2014 [32]    | Total effective rate of clinical efficacy | OR 2.80 [0.50, 15.73]     | No adverse reaction |
|                          | Improvement rate of cough               |                            |                |
|                          | Improvement rate of throat itching      |                            |                |
|                          | Improvement rate of cough up phlegm     |                            |                |
|                          | Total effective rate of airway responsiveness | OR 4.26 [0.81, 22.53]   |                |
| Li 2015 [33]             | Total effective rate of clinical efficacy | OR 1.78 [0.39, 8.09]     | No adverse reaction |
|                          | Recurrence rate                         | 0.16 [0.05, 0.47]         |                |
| Wang and Xie 2015 [34]   | Total effective rate of clinical efficacy | OR 2.22 [0.62, 7.97]     | No adverse reaction |
|                          | Recurrence rate                         |                            |                |
| Wei and Qin 2019 [35]    | Total effective rate of clinical efficacy | OR 5.57 [0.62, 50.03]    | No adverse reaction |
|                          | Total effective rate of airway responsiveness | OR 8.27 [0.97, 70.73]  |                |
| Wang 2020 [16]           | Total effective rate of clinical efficacy | OR 7.21 [1.48, 35.07]   | Not reported   |

RCT: randomized controlled trial, TNF: tumor necrosis factor, IL: interleukin, FEF50: forced expiratory flow at 50% of forced vital capacity, FEF75: forced expiratory flow at 75% of forced vital capacity, MMEF75/25: maximal mid-expiratory flow, OR: odds ratio, and MD: mean difference.
of blindness and the ability to ensure the presence of selective reports; two studies had incomplete outcome data. Despite the low-quality research methods, the careful evaluation of literature can be used to compensate for the factivity and creditability of results. Lastly, besides the different dosages of HQXXD compositions and the drugs in the control group, the methods, as well as duration of treatment, were not uniform. The presence of these biases might skew the research’s results. However, our research primarily focused on the use of HQXXD, particularly RA, HRA, HS, and RS for individuals with CVA, so there is no special regulation on the dose. The baselines for inclusion in the literature are not considerably different, and the studies that were included were RCTs with consistent diagnostic criteria. Based on interventions, controls, and duration of treatment, we assessed the heterogeneity as low. Based on methodological quality, statistical model, and sample size, the sensitivity analysis demonstrated that the robustness and reliability of the pooled results were fair. It was evident from the funnel plot that there is no remarkable publishing bias.
Furthermore, for total effective rates of clinical efficacy and airway responsiveness outcomes, the GRADE levels of evidence were moderate; for recurrence rate outcomes, the GRADE level of evidence is low. The high risk of bias was the main reason for a lower level; high-quality and multicenter RCTs are required to produce better evidence.

5. Conclusion

In brief, HQXXD is safe and effective in the treatment of CVA patients, which benefits clinical efficacy and airway responsiveness, reduces the recurrence rate, and has no adverse effects. More high-quality, multicenter, large-sample RCTs are required for the purpose of gathering better evidence because of the potential risk of bias. This systematic review offers more evidence regarding the effectiveness and safety of HQXXD for CVA and helps clinicians in decision-making when treating CVA patients. In addition, it can provide a basis for future basic and clinical research.

Data Availability

The extracted data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Authors’ Contributions

Cong Wang and Qingqing Xia contributed equally to this work. Cong Wang, Qingqing Xia, Weilong Jiang, and Huizhe Zhang conceived and designed the systematic review and wrote the manuscript. Cong Wang, Qingqing Xia, and Beina Hu conducted the database search, assessed studies for inclusion, and extracted and analyzed the data. Qingqing Xia performed supervision and project administration. Huizhe Zhang analyzed the data, arbitrated any disagreements, performed supervision, and revised the manuscript. All authors read and approved the final manuscript.

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Supplementary Materials

Supplementary File 1. PRISMA 2020 checklist. Supplementary File 2. Search strategy in PubMed for example. Table S1. Risk of bias of included RCTs. (Supplementary Materials)

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