REVIEW

3725

Systematic review and meta-analysis of randomized controlled trials on Wenxin keli

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Objective: The aim of the study was to evaluate the effectiveness, safety, and cost associated with Wenxin keli in the treatment of cardiovascular diseases based on meta-analysis.

Methods: The terms "Wenxin keli" and "Wenxin" were used as the search terms in the PubMed, ProQuest, Springer, the Cochrane Library, CNKI (China National Knowledge Infrastructure), VIP (Chinese Scientific Journals Database), and Wan fang electronic databases (from January 2000 to October 2015). Relevant print journals and conference papers were also searched. Studies on randomized controlled trials (RCTs) of Wenxin keli used in the treatment of cardiovascular diseases were screened, and its indications were classified. Meta-analysis of these studies was conducted using the RevMan 5.2 software.

Results: A total of 49 RCTs (n=4,610) were included, 29 of which focused on arrhythmia, seven on angina, seven on heart failure, two on viral myocarditis, and four on menopausal syndrome. Analysis of the therapeutic indications of Wenxin keli showed that it was comparatively more curative and effective than other available treatments for cardiovascular diseases.

Conclusion: Wenxin keli showed better clinical efficacy in the treatment of arrhythmia, angina, and heart failure; however, more high-quality evidence is needed to support its use in the clinical setting.

Keywords: Wenxin keli, cardiovascular disease, meta-analysis, systematic review

Introduction

The number of patients affected by cardiovascular disease is steadily increasing because of socioeconomic development and modern lifestyles.¹ A report from the World Health Organization reveals that chronic noncommunicable diseases have now become the leading cause of death worldwide. In 2008, 36 million individuals died of chronic noncommunicable diseases (63% of total deaths), of which 48% died of cardiovascular diseases.² In recent years, there has been a shift in the medical paradigm, and cardiovascular and cerebrovascular diseases have now become a serious threat to public health.³ According to a report on Cardiovascular Diseases in China (2013), presented at the China Heart Congress, about one-fifth of all Chinese adults are currently affected with cardiovascular and cerebrovascular diseases ranks first among all causes of death.⁴ Social and economic development has led to dramatic changes in lifestyles, including an increase in energy intake, less manual labor, accelerated pace of life, competitive pressures, and exposure to other risk factors associated with cardiovascular and other chronic diseases, which are causes for concern.⁵

Wenxin keli consists of several kinds of Chinese herbs including Huang Jing, *Codonopsis*, amber, *Panax*, and nard. Wenxin keli has been used clinically in the treatment of qi and yin deficiency, systolic blood stasis due to restless heart palpitations,

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shortness of breath, chest pain, premature ventricular contractions (PVC), and atrial premature beats. It is one of the main forms of treatment for cardiovascular disease in Chinese medicine. In recent years, many researchers have evaluated the use of Wenxin keli in the treatment of cardiovascular disease; however, further evaluation is necessary. Therefore, the present study aimed to conduct a comprehensive evaluation of the efficacy and safety of Wenxin keli, and provide the basis for its use as a medication for cardiovascular disease.

Materials and methods Literature search

We performed systematic searches for randomized controlled trials (RCTs) designed to evaluate the clinical efficacy of Wenxin keli in CNKI (China National Knowledge Infrastructure), Wan fang, VIP (Chinese Scientific Journals Database), PubMed, the Cochrane Library, Springer, and ProQuest from January 1, 2000 to September 7, 2015 using "Wenxin keli" and "Wenxin" as the search terms.

Inclusion and exclusion criteria

Based on the Cochrane Collaboration Handbook standards, the following inclusion criteria were formulated for the selected literature: all published domestic and international RCTs on Wenxin keli; comparable baseline test data; interventions with individual drugs and Wenxin keli doses of 9 g, three times/day; any particular course of treatment; publications in Chinese and English. Diagnostic criteria used in the present study were based on authoritative Chinese and other countries diagnostic criteria. The exclusion criteria were as follows: duplicate publications, reports of combination therapy effects on treatment, descriptive studies, studies involving animal testing, conflicting before and after data (such as, the sum of the data not matching the total), and reports without statistical indicators.

Quality assessment

Study quality was evaluated with an improved version of the Jadad questionnaire, considering mainly four aspects: 1) random sequence generation (2 points); 2) randomized hiding (2 points); 3) blinding (2 points); 4) a withdrawal period (1 point). Two reviewers independently completed the assessment, and the mean score of the two reviewers was used as the final quality score of the selected studies. In the assessment of RCTs, 1–3 points were considered as low quality, and 4–7 points indicated high quality.

A unified data extraction sheet was derived, based on blinding characteristics in previously published medical literature, for use by the two reviewers. The information thereby extracted was then cross-checked. The extracted data included:

- document specifications: first author, publication year, and title;
- subjects: disease, diagnostic criteria, inclusion and exclusion criteria, sample size, etc;
- interventions: medication, dosage, route of administration, duration of treatment, etc;
- 4) results: efficacy indicators.

Statistical analysis

The RevMan 5.2 software, provided by the Cochrane Collaboration, was used to conduct the meta-analysis. Count data were used to determine the odds ratio (OR) and 95% confidence interval (CI) for the efficacy analysis of effect size, whereas measurement data were used to determine the standardized mean difference. Heterogeneity of the included studies was expressed in terms of *P* and *P*². If *P*>0.1 and $P^2 < 50\%$, the result of the test for heterogeneity was considered not statistically significant, and the fixed effects model was used for meta-analysis. For contrast, the random effects model was also applied to the data when $P \leq 0.1$ and $P^2 \geq 50\%$.

Results

Retrieval results and quality assessment

We searched 2,970 potentially relevant articles in CNKI, 3,488 in Wan fang, 2,393 in VIP, and two each in PubMed, Springer, the Cochrane Library, and ProQuest. We retrieved 2,274 reports after reading the abstracts, and reports involving animal studies, pharmacological studies, and systems analyses were excluded. After screening the full texts of 663 documents that were selected following application of the inclusion and exclusion criteria, 29 studies on arrhythmia were included,⁶⁻³⁴ of which nine were on PVC, seven on angina,³⁵⁻⁴¹ seven on heart failure,⁴²⁻⁴⁸ two on viral disorders,^{49,50} and four on climacteric syndrome,⁵¹⁻⁵⁴ as shown in Figure 1 and Table 1.

The quality assessment of the studies was performed by two independent reviewers. Of the 49 studies included (Figure 1 and Table 1), only two studies were found to be of high quality (4 points).^{6–54} The results of specific assessment are presented in Table 1.

Results of meta-analysis

Meta-analysis of Wenxin keli in the treatment

of arrhythmia

Clinical efficacy

There were 29 reports on the use of Wenxin keli in the treatment of arrhythmia, including nine on PVC. The results showed that Wenxin keli exhibited better clinical efficacy in



Figure I Study selection steps.

Abbreviations: CNKI, China National Knowledge Infrastructure; VIP, Chinese Scientific Journals Database; WF, Wan fang.

Study	Indications	Number of cases	Treatment	Interventions		Outcomes	Jadad
		(test/control groups)	(days)	Drug test	Control drug		score
Gao ⁶	Arrhythmia	48/48	28	Wenxin keli	Propafenone	I, 4	2
Liu et al ⁷	Arrhythmia	75/75	28	Wenxin keli	Propafenone	I	2
Wang ⁸	Arrhythmia	60/60	28	Wenxin keli	Propafenone	I, 4	2
Cui ⁹	Arrhythmia	60/60	28	Wenxin keli	Propafenone	I, 2	2
Li et al ¹⁰	Arrhythmia	48/35	28	Wenxin keli	Propafenone	I, 2, 3	1
Xie ¹¹	Arrhythmia	34/32	28	Wenxin keli	Propafenone	I, 4	2
Zou and Zhao ¹²	Arrhythmia	102/101	28	Wenxin keli	Propafenone	I	2
Wang ¹³	Arrhythmia	75/75	28	Wenxin keli	Propafenone	I, 4	2
Lou ¹⁴	Arrhythmia	53/50	28	Wenxin keli	Propafenone	I	2
Shi¹⁵	Arrhythmia	32/30	28	Wenxin keli	Propafenone	I, 4	2
Wang ¹⁶	Arrhythmia	58/64	28	Wenxin keli	Propafenone	2, 4	2
Jin and Huang ¹⁷	Arrhythmia	20/20	28	Wenxin keli	Propafenone	3	3
Xue ¹⁸	Arrhythmia	126/72	28	Wenxin keli	Propafenone	3	3
Ren and Qiao ¹⁹	Arrhythmia	43/21	28	Wenxin keli	Propafenone	3	3
Wu and Yue ²⁰	Arrhythmia	48/33	28	Wenxin keli	Propafenone	3	2
Li and Shen ²¹	Arrhythmia	40/37	28	Wenxin keli	Amiodarone	I, 4	2
Wang ²²	Arrhythmia	46/30	28	Wenxin keli	Amiodarone	I, 4	2
Pang ²³	Arrhythmia	56/58	28	Wenxin keli	Amiodarone	I, 4	2
Xu et al ²⁴	Arrhythmia	68/61	28	Wenxin keli	Amiodarone	I, 4	2

Table I Basic characteristics of included studies

(Continued)

Table I (Continued)

Study	Indications	Number of cases Treatment Interventions			Outcomes		
occup	marcacions	(test/control groups)	(days)	Drug test	Control drug	euccomes	score
V:- 25	A		20	Manuin hali	Auria deura una	1.4	2
X1a ²⁵	Arrnythmia	50/50 12/20	28		Amiodarone	1,4	2
Sun \\\27	FVC	52/31	20	Wenxin keli	Propatenone	1, 2, 3	2
VVU Mang ²⁸	PVC	5 1 /35	20	Wonyin keli	Propatenone	1, 2	2
Guo ²⁹	PVC	52/52	20	Wonyin keli	Propatenone	1 2 2	2
lin ³⁰	PVC	22/22	20	Wonyin koli	Propatenone	1, 2, 3	2
Zhang of al ³	PVC	39/39	20	Wanyin kali	Propatenone	1, 2, 5	2
	PVC	37/30	20	Wanyin kali	Propatenone	2, 5	2
Wang ³³	PVC	60/60	20	Wenzin keli	Propafenone	2 3	2
Yan ³⁴	PVC	60/60	28	Wenxin keli	Propafenone	2, 3	2
Notes: I, clinical	efficacy; 2, ECG e	efficacy; 3, clinical symptoms o	of heart palpitatio	ns, shortness of breath, dizziness	, insomnia, chest tightness, etc; 4	4, adverse reacti	ons.
Vu et el35	Angina	40/25		Conventional treatment	Conventional treatment	1.2	- <u>-</u> -
Tu et als	Angina	40/35	28	Wenxin keli	Conventional treatment	1, 2	Z
Shu and Li ³⁶	Angina	37/37	28	Conventional treatment +	Conventional treatment	1	2
				Wenxin keli			
Ye et al ³⁷	Angina	36/36	28	Conventional treatment +	Conventional treatment	12	2
i e et al	7 115114	50,50	20	Wonyin koli	Conventional d'eathene	·, <u>-</u>	-
14/2: 24 2138	A	50/50	20			1.2	2
vvei et also	Angina	50/50	28	Conventional treatment +	Conventional treatment	Ι, Ζ	2
				Wenxin keli			
Yuan ³⁹	Unstable	47/47	28	Conventional treatment +	Conventional treatment	I, 4	2
	angina			Wenxin keli			
Wei and	Unstable	50/50	28	Conventional treatment +	Conventional treatment	Ι, 3	2
Deng ⁴⁰	angina			Wenxin keli			
Yuan and Wei ⁴¹	Unstable	80/80	28	Conventional treatment +	Conventional treatment	1.3.4	2
	angina			Wenxin keli		, ,	
		(C) D () () () ()					
Notes: I, angina p	bectoris; 2, ECG	efficacy; 3, changes in the indic	cator lipids; 4, ad	verse reactions.			
Yu et al ⁴²	Chronic	37/35	56	Conventional treatment +	Conventional treatment	1, 2, 3	4
	heart failure			Wenxin keli			
Yang and	Chronic	40/40	56	Conventional treatment +	Conventional treatment	1. 3. 4	2
Dong ⁴³	heart failure			Wenzin keli		., ., .	
Kong of al ⁴⁴	Chronic	20/20	54	Conventional treatment +	Conventional treatment	246	C
Kong et al		30/30	30		Conventional treatment	2, 7, 0	2
24	neart failure	27/25	- /			-	
Xu⁴⁵	Chronic	37/35	56	Conventional treatment +	Conventional treatment	3	4
	heart failure			Wenxin keli			
Hu ⁴⁶	Chronic	50/48	56	Conventional treatment +	Conventional treatment	4	2
	heart failure			Wenxin keli			
Yu ⁴⁷	Congestive	35/37	56	Conventional treatment +	Conventional treatment	1, 2, 3, 5	2
	heart failure			Wenxin keli			
Wang ⁴⁸		35/35	56	Conventional treatment +	Conventional treatment	12467	2
t t ang	beent failune	55,55	50		Conventional d'eathene	1, 2, 1, 0, 7	-
Notos: L clinical a	ficacy: 2 TCM of	undromos: 2 clinical ochocardi	ography: 1 plasm		ninuto walking distance measurem	ont: 7 advorce	mastions
			ogi apity, ¬, plasifi	a Divi values, J, near t late, 0, 0-1	minute maining distance measurem		
Yang ⁴⁹	Children	34/34	14	Conventional treatment +	Conventional treatment	1, 2	2
	with viral			Wenxin keli			
	myocarditis						
Deng ⁵⁰	Children	30/28	14	Conventional treatment +	Conventional treatment	1	2
- 0	with viral			Wenxin keli			
	myocarditis			t tenxin ken			
Notes: L clinical	efficacy: 2 creati	ne kinase (CK-MB) change: 3	adverse reaction	2			
	enicacy, 2, creati	The Kinase (CK-ITB) change, 5,		s.			
Liu and Ren⁵	Climacteric	60/58	84	Conventional treatment +	Conventional treatment +	I	2
	syndrome			Wenxin keli	metoprolol		
Lei ⁵²	Climacteric	42/38	84	Conventional treatment +	Conventional treatment +	I	2
	syndrome			Wenxin keli	metoprolol		
Li and Miao53	Climacteric	33/33	60	Wenxin keli	Oryzanol + propranolol +	1	2
	syndrome				vitamin B complex		
Hu ⁵⁴	Climacteric	25/23	60	Wenxin keli	Oryzanol + propranolol +	1	2
. 14	syndromo	_3, _3			vitamin B complex		-
	syndrome				vitamin & complex		

Notes: I, clinical efficacy; 2, adverse reactions.

Abbreviations: BNP, brain natriuretic peptide; CK-MB, creatine kinase MB isoenzyme; ECG, electrocardiogram; PVC, premature ventricular contractions; TCM, traditional Chinese medicine.

the treatment of arrhythmia (OR =1.74, 95% CI [1.28, 2.35], P=0.0003; Figure 2) compared to propafenone. Also, in comparison to amiodarone, Wenxin keli again exhibited better clinical efficacy in the treatment of arrhythmia (OR =2.28, 95% CI [1.33, 3.89], P=0.003; Figure 3).

Five studies considered the use of Wenxin keli in the treatment of PVC. The meta-analysis showed that it exhibited better clinical efficacy than propafenone (OR =2.92, 95% CI [1.72, 4.96], P<0.0001; Figure 4).

Efficacy of Wenxin keli on electrocardiogram

Three studies reported on the efficacy of Wenxin keli on electrocardiogram (ECG). Meta-analysis of the random effects model showed no significant difference between Wenxin keli and propafenone in the treatment of arrhythmia based on the ECG (OR =2.15, 95% CI [0.58, 7.97], P=0.25; Figure 5).

Eight studies reported on the use of Wenxin keli in the treatment of PVC. The meta-analysis showed that Wenxin keli showed better efficacy than propafenone based on the ECG (OR =2.19, 95% CI [1.45, 3.30], P=0.0002; Figure 6).

Secondary outcomes

Five studies reported on the effect of Wenxin keli treatment on secondary efficacy variables in PVC. Heterogeneity was minimal; thus, the fixed effects model was applied to the study that reported on secondary efficacy variables, in addition to dizziness. The results showed that Wenxin keli exhibited better efficacy, in addition to dizziness (Table 2).

Adverse reactions

A total of eleven studies reported adverse reactions in the treatment of arrhythmia. Wenxin keli showed a lower incidence of adverse reactions, with reports of mild adverse reactions and favorable clinical application and safety, in comparison to both propafenone and amiodarone (Figures 7 and 8).

Six studies reported on adverse reactions in the treatment of PVC. Meta-analysis of the fixed effects model showed that Wenxin keli exhibited a lower incidence of adverse reactions (OR =0.32, 95% CI [0.16, 0.64], *P*=0.001; Figure 9) compared to amiodarone.

Publication bias

A funnel plot (Figure 10) showed that the studies reporting the use of Wenxin keli in the treatment of arrhythmia, included in the analysis, were substantially symmetric. This suggests less publication bias in these reports. According to the Cochrane Handbook, funnel plot analysis should not be performed for other indications if there are less than ten studies.

Meta-analysis of Wenxin keli in the treatment of angina

Angina pectoris

Six reports, included in the analysis, reported on the use of Wenxin keli in the treatment of angina pectoris, of which three specifically focused on angina and three on unstable angina. Meta-analysis of the fixed effects model showed that the clinical efficacy of Wenxin keli combined with conventional therapy in the treatment of angina was significantly better than conventional therapy alone (OR =3.12, 95% CI [1.77, 5.52], P<0.0001; Figure 11). The clinical efficacy of Wenxin keli combined with conventional therapy in the treatment of unstable angina was also significantly better than conventional therapy alone (OR =3.97, 95% CI [1.92, 8.22], P=0.0002; Figure 12).

Study or subgroup	Experime Events	ental Total	Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95% Cl		Odds ratio M–H, fixed	, 95% CI	
Cui ⁹	44	60	44	60	18.1	1.00 (0.45, 2.25)				
Gao ⁶	41	48	41	48	9.2	1.00 (0.32, 3.11)				
Li et al ¹⁰	42	48	26	35	5.8	2.42 (0.77, 7.60)		+		
Liu et al ⁷	62	75	48	75	12.9	2.68 (1.25, 5.74)				
Lou ¹⁴	46	53	42	50	8.8	1.25 (0.42, 3.75)				
Shi ¹⁵	30	32	25	30	2.5	3.00 (0.54, 16.81)		-		
Wang ¹³	69	75	57	75	7.1	3.63 (1.35, 9.76)				
Wang ⁸	51	60	51	60	11.8	1.00 (0.37, 2.72)				
Xie ¹¹	30	34	20	32	3.7	4.50 (1.27, 15.95)				
Zou and Zhao ¹²	86	102	82	101	20.0	1.25 (0.60, 2.59)		-		
Total (95% CI)		587		566	100	1.74 (1.28, 2.35)			•	
Total events	501		436						•	
Heterogeneity: χ^2 =11.28, <i>df</i> =9 (<i>P</i> =0.26); Test for overall effect: <i>Z</i> =3.58 (<i>P</i> =0.0003)		; /²=20% 3)				0.01 Favors (0.1 1 experimental)	10 Favors (cont	100 trol)	

Figure 2 Meta-analysis of Wenxin keli and propafenone in the treatment of arrhythmia. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.

Study or Experimental subgroup Events Total		Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95% 0		Odo M–ł	ls ratio I, fixed, 95	% CI		
Li and Shen ²¹	34	40	31	37	26.4	1.10 (0.32, 3.76)	-	5. S. S.			
Pang ²³	53	56	47	58	13.5	4.13 (1.09, 15.72)			-		
Wang ²²	38	46	23	30	26.4	1.45 (0.46, 4.52)				_	
Xia ²⁵	48	50	43	50	9.4	3.91 (0.77, 19.83)			-		
Xu et al ²⁴	62	68	48	61	24.4	2.80 (0.99, 7.90)			-	<u> </u>	
Total (95% CI)		260		236	100	2.28 (1.33, 3.89)				•	
Total events	235		192						-		
Heterogeneity: 2	2=3.30, df=4	4 (<i>P</i> =0.51);	/2=0%						-		
Test for overall e	effect: 7 =3.0	1 (P=0 003)				0.01	0.1	1	10	100
		. (. 0.000	/				Favors	(experime	ntal) Fa	avors (cont	rol)

Figure 3 Meta-analysis of Wenxin keli and amiodarone in the treatment of arrhythmia. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.

Study orExperimentalsubgroupEventsTotal		Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95% Cl		Odds M–H	s ratio , fixed,	95% CI		
Wu ²⁷	51	54	23	31	10.1	5.91 (1.44, 24.35)		5765 67o			
Wang ²⁸	53	60	21	30	20.4	3.24 (1.07, 9.84)			-		
Sun ²⁶	29	32	23	31	13.7	3.36 (0.80, 14.13)			+		
Jin ³⁰	53	60	21	30	20.4	3.24 (1.07, 9.84)				_	
Guo ²⁹	46	53	43	53	35.4	1.53 (0.53, 4.37)			+		
Total (95% CI)		259		175	100	2.92 (1.72, 4.96)				+	
Total events	232		131								
Heterogeneity:	$\chi^2 = 2.52, df = 4$	(P=0.64);	/2=0%			I		- 1	-		
Test for overall	effect: Z=3.97	(P<0.000	1)			0.0	01	0.1	1	10	100
		,	,			Fa	avors	(experimer	ntal)	Favors (cont	rol)

Figure 4 Meta-analysis of Wenxin keli and propafenone in the treatment of PVC. Abbreviations: CI, confidence interval; PVC, premature ventricular contractions; M–H, Mantel–Haenszel.

Study or Experimental subgroup Events Total		Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95% 0	Odds ratio CI M–H, fixed, 95% CI					
Cui ⁹	54	60	39	60	33.7	4.85 (1.79, 13.13)					
Li et al ¹⁰	43	48	25	35	31.2	3.44 (1.06, 11.21)			H		
Wang ¹⁶	44	58	53	64	35.1	0.65 (0.27, 1.58)		-	•		
Total (95% CI)		166		159	100	2.15 (0.58, 7.97)					
Total events	141		117						- 1		
Heterogeneity: τ^2	=1.07; χ ² =9	.97, df=2 (l	P=0.007); /2=	80%					-		
Test for overall ef	fect: Z=1.15	(P=0.25)					0.01	0.1	1	10	100
		(Favors	(experimen	ntal)	Favors (contr	ol)

Figure 5 Meta-analysis of Wenxin keli and propafenone in the treatment of arrhythmia. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.

Study or subgroup	Experime Events	ental Total	Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95% C	I	Odds ratio M–H, fixed) I, 95% Cl	
Guo ²⁹	60	65	62	65	15.2	0.58 (0.13, 2.54)				
Jin ³⁰	53	60	24	30	11.9	1.89 (0.57, 6.24)			•	
Li ³²	28	32	24	32	9.6	2.33 (0.62, 8.72)		+		
Sun ²⁶	23	32	22	31	20.0	1.05 (0.35, 3.12)				
Wang ³³	58	60	44	60	4.7	10.55 (2.30, 48.28)				_
Wu ²⁷	48	54	25	35	10.7	3.20 (1.04, 9.82)		-		
Yan ³⁴	55	60	45	60	12.0	3.67 (1.24, 10.86)			-	
Zhang et al ³¹	33	39	32	38	15.9	1.03 (0.30, 3.53)		-		
Total (95% CI)		402		351	100	2.19 (1.45, 3.30)			•	
Total events	358		278						0.500	
Heterogeneity: 2	2=11.77, df=	7 (<i>P</i> =0.11); /²=41%				<u> </u>	- + - +		
Test for overall e	effect: Z=3.73	(P=0.000	(2)				0.01	0.1 1	10	100
		. 5.000	_,				Favors	(experimental)	Favors (cont	trol)

Figure 6 Meta-analysis of Wenxin keli and propafenone in the treatment of PVC.

Abbreviations: CI, confidence interval; PVC, premature ventricular contractions; M–H, Mantel–Haenszel.

Table 2 Meta-analysis c	of secondary	^r efficacy	[,] variables ir	n treatment	of arrh	ythmia
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Outcomes	Number of	Number	Hetero	ogeneity	Model	Meta-analysis			
	included studies	of cases	1 ²	P-value		OR (95% CI)	P-value		
Palpitations	4	383	0%	0.99	Fixed	3.29 (1.64, 6.61)	0.0008		
Chest tightness	5	451	0%	0.70	Fixed	3.61 (2.22, 5.87)	<0.00001		
Restless sleep	4	373	0%	0.50	Fixed	2.49 (1.40, 4.43)	0.002		
Dizziness	4	379	55%	0.09	Random	2.53 (0.87, 7.35)	0.09		
Shortness of breath	5	462	0%	0.94	Fixed	3.00 (1.74, 5.19)	<0.0001		

Abbreviations: CI, confidence interval; OR, odds ratio.

Study or Experimental subgroup Events Total		Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95% 0		Odds M–H,	ratio fixed,	95% CI		
Gao ⁶	2	52	10	64	19.9	0.22 (0.05, 1.03)				Annas 200	
Shi ¹⁵	0	32	3	30	8.2	0.12 (0.01, 2.44)	+		_		
Wang ¹⁶	2	58	10	64	21.2	0.19 (0.04, 0.92)			_		
Wang ¹³	6	75	16	75	34.0	0.32 (0.12, 0.87)			_		
Wang ⁸	2	60	7	60	15.6	0.26 (0.05, 1.31)			-		
Xie ¹¹	3	34	0	32	1.1	7.22 (0.36, 145.56)	-			→
Total (95% CI)		311		325	100	0.32 (0.18, 0.58)		•	-		
Total events	15		46								
Heterogeneity:	γ ² =5.25. df=5	(P=0.39);	/²=5%						-		
Test for overall	effect: 7=3 76	(P=0 000	2)				0.01	0.1	1	10	100
	5 D		-,				Favors	(experimen	tal)	Favors (cont	rol)

Figure 7 Meta-analysis of Wenxin keli, propafenone, and associated adverse reactions on arrhythmia. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.

Study or Experimental subgroup Events Tota		ental Total	Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95%	CI	Oddsr M–H, fi	atio xed, 9	95% CI	
Li and Shen ²¹	10	40	17	37	17.1	0.39 (0.15, 1.03)				w 197 201	
Pang ²³	3	56	18	58	21.6	0.13 (0.03, 0.46)					
Wang ²²	1	46	9	30	13.7	0.05 (0.01, 0.44)	+				
Xia ²⁵	3	50	15	50	18.2	0.15 (0.04, 0.55)					
Xu et al ²⁴	4	68	23	61	29.4	0.10 (0.03, 0.32)					
Total (95% CI)		260		236	100	0.16 (0.09, 0.27)		•			
Total events	21		82					-			
Heterogeneity:	2=5.12, df=4	(P=0.27);	l ² =22%						-		
Test for overall e	effect: Z=6.72	(P<0.000	01)				0.01	0.1	1	10	100
							Favors	s (experimenta	I)	Favors (conti	rol)

Figure 8 Meta-analysis of Wenxin keli, amiodarone, and associated adverse reactions on arrhythmia. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.

Study or Experimental subgroup Events Total		Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95% (CI	Odds M–H,	ratio fixed, 9	5% CI		
Guo ²⁹	0	65	3	65	11.5	0.14 (0.01, 2.69)	+		-		
Jin ³⁰	3	60	1	30	4.2	1.53 (0.15, 15.33)					
Sun ²⁶	2	32	4	31	12.6	0.45 (0.08, 2.66)			-	-	
Wang ³³	4	60	8	60	24.8	0.46 (0.13, 1.63)		_	-		
Yan ³⁴	0	60	6	60	21.4	0.07 (0.00, 1.26)	+	-	-		
Zhang et al ³¹	2	39	8	38	25.5	0.20 (0.04, 1.03)			-		
Total (95% CI)		316		284	100	0.32 (0.16, 0.64)		-	•		
Total events	11		30								
Heterogeneity:	γ ² =3.94, df=5	5 (P=0.56);	/2=0%						-		
Test for overall e	effect: 7=3 19	Q (P=0 001)				0.01	0.1	1	10	100
	5.1000. <u>2</u> -0.10	0.001	/				Favors	s (experiment	al) F	avors (conti	rol)

Figure 9 Meta-analysis of Wenxin keli, propafenone, and associated adverse reactions on PVC. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.



Figure 10 Funnel plot of Wenxin keli and propafenone in the treatment of arrhythmia. Abbreviations: SE. standard error of the mean: OR. odds ratio.

ECG efficacy

Two studies reported on the efficacy of Wenxin keli in the treatment of angina based on the ECG. Meta-analysis of the fixed effects model showed that the efficacy of Wenxin keli combined with conventional therapy was not significantly different from that of the conventional therapy alone (OR =2.02, 95% CI [0.65, 6.24], P=0.22; Figure 13).

Adverse reactions

Three studies mentioned the development of adverse reactions to Wenxin keli in the treatment of angina, two of which could not be compared because the experimental and control groups were not described separately. The latter test group and six cases (15%) in the control group exhibited no adverse reactions.

Meta-analysis of Wenxin keli in the treatment of heart failure

Clinical efficacy

Two studies reported on the clinical efficacy of Wenxin keli in the treatment of chronic heart failure. Meta-analysis showed that Wenxin keli combined with conventional treatment showed no greater clinical efficacy (OR =2.62, 95% CI [0.91, 7.56], P=0.07; Figure 14) compared to the conventional treatment group.

Secondary efficacy variables

Analysis of secondary efficacy end points showed that Wenxin keli combined with conventional treatment showed better efficacy in left ventricular ejection fraction (LVEF) values, plasma brain natriuretic peptide (BNP) levels, and stroke volume (Table 3).

Meta-analysis of Wenxin keli in the treatment of viral infections

Clinical efficacy

Two studies reported on the clinical efficacy of Wenxin keli in the treatment of viral infections. Meta-analysis showed that Wenxin keli combined with conventional treatment exhibited better clinical efficacy (OR =4.89, 95% CI [1.30, 18.38], P=0.02; Figure 15) compared to conventional treatment.

Adverse reactions

Two studies investigated adverse reactions associated with Wenxin keli in the treatment of viral infections. No adverse reactions were reported in either study, suggesting the safety of Wenxin keli.

Meta-analysis of Wenxin keli in the treatment of climacteric syndrome

Clinical efficacy

Four studies reported on the clinical efficacy of Wenxin keli in the treatment of climacteric syndrome, of which two compared Wenxin keli combined with conventional therapy to conventional therapy alone, and the other two compared Wenxin keli to a combination of oryzanol, propranolol, and vitamin B complex. Meta-analysis showed that in comparison to conventional treatment alone, Wenxin keli combined



Figure II Meta-analysis of Wenxin keli and propafenone in the treatment of angina. Abbreviations: Cl, confidence interval; M–H, Mantel–Haenszel.

Study or subgroup	Experim Events	ental Total	ntal Control Weight Odds ratio Total Events Total (%) M–H, fixed, 95% Cl				Odds ratio ₀ Cl M–H, fixed, 95% Cl				
Wei et al ³⁸	45	50	38	50	47.0	2.84 (0.92, 8.79)					
Yuan and Wei41	27	30	23	30	28.5	2.74 (0.63, 11.82)	-	_			
Yuan ³⁹	44	47	31	47	24.5	7.57 (2.03, 28.22)					
Total (95% CI)		127		127	100	3.97 (1.92, 8.22)		+			
Total events	116		92								
Heterogeneity: χ^2	e=1.51, df=2	(P=0.47);	/2=0%			F					
Test for overall of	fect: 7=3 72	(P=0,000)	0.0	1 0.1 1	10	100					
	1001. 2-0.72	(i =0.000)	-)			Fa	vors (experimental)	Favors (cont	rol)		

Figure 12 Meta-analysis of Wenxin keli and propafenone in the treatment of unstable angina. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.

Study or subgroup	Experimental Events Total		Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95% (CI	Odd M–H	s ratio , fixed	, 95% CI	
Wei et al ³⁸	39	50	25	50	51.1	3.55 (1.49, 8.45)					
Ye et al37	21	36	20	36	48.9	1.12 (0.44, 2.85)			-	<u> </u>	
Total (95% CI)		86		86	100	2.02 (0.65, 6.24)				•	
Total events	60		45								
Heterogeneity: $r^2=0.45$, $\chi^2=3.14$, $df=1$ (<i>P</i> =0.08); $l^2=68\%$							0.01	0.1	1	10	 100
					Favors			Favors (cont	rol)		

Figure 13 Meta-analysis of Wenxin keli combined with conventional therapy in the treatment of angina. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.

Study or subgroup	Experime Events	ental Total	Control Events	Total	Weight (%)	Odds ratio M–H, fixed, 95% (Odds rati M–H, fixe	o d, 95% Cl	
Yang and Dong43	37	37	33	35	10.0	5.60 (0.26, 120.80)			
Yu et al ⁴²	55	60	48	58	90.0	2.29 (0.73, 7.17)		-		
Total (95% CI)		97		93	100	2.62 (0.91, 7.56)			-	
Total events	92		81							
Heterogeneity: $\chi^2 =$	0.29, <i>df</i> =1 (P=0.59); I	² =0%							
Test for overall effect: Z=1.78 (P=0.07)							0.01	0.1	1 10	100
,							Favors	(experimental)	Favors (Control)	

Figure 14 Meta-analysis of Wenxin keli combined with conventional treatment for chronic heart failure. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.

Outcomes	Number of	Number	Hetero	ogeneity	Model	Meta-analysis		
	included studies	of cases	1 ²	P-value		MD (95% CI)	P-value	
LVEF values	4	310	57%	0.08	Random	0.76 (0.41, 1.12)	<0.0001	
Plasma BNP levels	3	238	99 %	<0.00001	Random	-5.92 (-9.70, -2.14)	0.002	
Stroke volume	3	212	29%	0.24	Fixed	0.50 (0.18, 0.83)	0.002	

Abbreviations: BNP, brain natriuretic peptide; CI, confidence interval; LVEF, left ventricular ejection fraction; MD, mean difference.

Study or subgroup	Experimental Events Total		Control Events	Total	Weight (%)	Weight Odds ratio (%) M–H, fixed, 95% C		Odds ratio I M–H, fixed, 95% Cl			
Deng⁵⁰	29	30	24	28	35.1	4.83 (0.51, 46.18)		1000			
Yang ⁴⁹	32	34	26	34	64.9	4.92 (0.96, 25.22)			-		
Total (95% CI)		64		62	100	4.89 (1.30, 18.38)			-		
Total events	61		50								
Heterogeneity: χ	² =0.00, <i>df</i> =1	(P=0.99);	/ ² =0%						+ +		
Test for overall e	ffect: Z=2.35	(P=0.02)					0.01	0.1	1 10	100	
							Favors	(experimenta	l) Favors (cont	Favors (control)	

Figure 15 Meta-analysis of Wenxin keli combined with conventional treatment for viral infections. Abbreviations: CI, confidence interval; M–H, Mantel–Haenszel.

Study or subgroup	Experimental Events Total		Control Events	Total	Weight al (%)	Odds ratio M–H, fixed, 95% 0	;	Odds ratio M–H, fixed, 95% Cl			
Lei ⁵²	36	42	22	38	35.7	4.36 (1.49, 12.82)			· · · · · · · · · · · · · · · · · · ·		
Liu and Ren ⁵¹	50	60	35	58	64.3	3.29 (1.39, 7.76)			-		
Total (95% CI)		102		96	100	3.67 (1.88, 7.18)			+		
Total events	86		57						6380		
Heterogeneity: χ	² =0.16, <i>df</i> =1	(P=0.69);	I2=0%						+ +		
Test for overall effect: $Z=3.80$ ($P=0.0001$)							0.01	0.1	1 10	100	
							Favors	s (experimenta	I) Favors (con	Favors (control)	

Figure 16 Meta-analysis of Wenxin keli combined with conventional treatment for climacteric syndrome. Abbreviations: Cl, confidence interval; M–H, Mantel–Haenszel.

with conventional treatment exhibited better clinical efficacy in the treatment of climacteric syndrome (OR =3.67, 95% CI 1.88, 7.18, *P*=0.0001; Figure 16). In comparison to the control drug (propranolol + oryzanol + vitamin B), Wenxin keli showed better clinical efficacy (OR =7.82, 95% CI [2.92, 20.95], *P*<0.0001; Figure 17).

Adverse reactions

No adverse reactions were reported in the literature regarding the use of Wenxin keli in the treatment of climacteric syndrome. Thus, comparisons between any control and corresponding experimental groups were not possible.

Discussion

Wenxin keli consists primarily of *Codonopsis*, Huang Jing, *Panax*, amber, nard, and other traditional Chinese herbs. It represents the first broad-spectrum treatment capable of affecting multiple ion channels (Na⁺, K⁺, and Ca²⁺) that can also significantly improve heart function (without causing arrhythmias), heart palpitations, chest tightness, and other associated symptoms. Modern pharmacological studies have confirmed that *Codonopsis* contains inulin and amino acids, and that it exerts anti-platelet aggregation, enhances immunity, and improves myocardial contractile effects.⁵⁵ Huang Jing exhibits lipid-lowering and anti-atherosclerotic effect, reduces blood pressure, and increases coronary blood flow.⁵⁶ *Panax* can increase coronary blood flow, inhibit self-discipline of the ectopic pacemaker sinus

node, reduce myocardial oxygen consumption, improve microcirculation, and regulate myocardial ischemia and hypoxia.⁵⁶ Nard plays a role in relieving depression, and pharmacological experiments show that it contains valerian ketones. These compounds can combine with specific proteins via ion channels in the myocardial cell membrane to reduce myocardial cell automaticity, extend the atrial action potential of ventricular muscle and conduction system time, interrupt reentry, and eliminate arrhythmias.⁵⁷ The therapeutic index of Wenxin keli on the heart, kidneys, and liver was within normal limits. Wenxin keli can enhance immune function, without causing significant adverse reactions, and shows no evidence of the side effects of myocardial ischemia and arrhythmia. It is therefore considered to be safe and effective.³

Limitations

Among the studies included in the meta-analysis, only two of them that had higher scores on the Jadad questionnaire were used to evaluate the quality, thereby potentially affecting the strength of the results. There were fewer documents available for some of the indications analyzed, and some studies had smaller sample sizes. Both of these factors represent limitations of the present study. No standards for RCTs have been published in People's Republic of China; therefore, allocation concealment and blinding were rarely mentioned in the studies included in the analysis. Thus, it is possible that most of the original



Figure 17 Meta-analysis of Wenxin keli in the treatment of climacteric syndrome. Abbreviations: Cl, confidence interval; M–H, Mantel–Haenszel. reports were inconclusive and the results were of low quality, thereby increasing bias. To address the aforementioned limitations and verify the results of the present study, additional high-quality RCT studies that employ larger sample sizes are required.

Conclusion

Based on the available evidence, meta-analysis is an effective method to prove the safety and efficacy of a particular treatment. The results of meta-analysis allow physicians and patients to choose the most effective treatment.

For systematic reviews of Wenxin keli in the treatment of cardiovascular disease, we used the indicators of angina pectoris total efficiency, ECG total efficiency, and adverse outcomes for comparison with the control groups that were treated with propafenone and amiodarone. The findings observed in the treatment of arrhythmia, PVC, angina pectoris, heart failure, viral myocarditis, and climacteric syndrome, among others, were derived from 49 studies. Overall, these studies reported favorable effects of Wenxin keli, regardless of whether it was used directly or as an adjuvant therapy. Furthermore, a low incidence of adverse reactions was evident among the studies analyzed.

Disclosure

The authors report no conflicts of interest in this work.

References

- Go AS, Mozaffarian D, Roger VL, et al. Heart disease and stroke statistics – 2014 update: a report from the American Heart Association. *Circulation*. 2014;129(3):399–410.
- World Health Organization. Global Status Report on Noncommunicable Diseases 2010. Geneva: WHO; 2011.
- 3. Wang J, Shui LM. 1991–2003 Ningbo City Jiangdong district residents cardiovascular disease mortality dynamic analysis. *Chin J Prevent Control Chronic Dis*. 2005;13(5):251–252.
- 4. Hu XL, Zhou JM, Chen XP. Role of AGXT2 in ADMA metabolism and the development of cardiovascular and cerebrovascular diseases. *Chin Pharmacol Bull*. 2015;05:601–605.
- Li YH, Zhou JM, Yang C, et al. Study on cardiovaseular and cerebrovaseular diseases and its influencing factors among 5 kinds of occupational populations in 6 provinces in China. *Chin J Health Education*. 2012;01:3–7.
- Gao P. Clinical analysis of 52 cases buchang Wenxin keli in treatment of arrhythmia. *Chin Comm Doc.* 2010;13:144.
- Liu JF, Cao PL, Ge YZ, et al. Curative effect analysis of Wenxin keli in treatment of cardiac arrhythmia. *Acta Acad Med Jiangxi*. 2009;09: 52–54.
- Wang YM. Clinical study of Wenxin keli in treatment of cardiac arrhythmia. *Chin Prac Med.* 2007;23:3–5.
- Cui YX. Clinical analysis of buchang Wenxin keli in treatment of arrhythmia. *Chin Prac Med.* 2010;26:165–166.
- Li YB, Sun CY, et al. Clinical observation of 48 cases on Wenxin keli in treatment of arrhythmia. *Chin Comm Doc.* 2006;22(09):43.
- Xie HQ. Clinical analysis of 34 cases on Wenxin keli in treatment of arrhythmia. J Kunming Med Coll. 2009;(3B):386–387.
- 12. Zou GX, Zhao BC. 102 cases of Wenxin keli in treatment of arrhythmia. *J Tradit Chin Med*. 2002;09:689–690.

- 13. Wang CY. 150 cases of Wenxin keli in treatment of arrhythmia. *Clin J Rational Drug Use.* 2012;28:65–66.
- Lou MP. 53 cases of Wenxin keli in treatment of arrhythmia. *Modern Medicine*. 2008;17:2627–2628.
- 15. Shi H. The efficacy of Wenxin keli in treatment of arrhythmia. *Chemists* (Academic Edition). 2012;10(12):174–175.
- Wang MS. Clinical observation on treatment of 58 cases with cardiac arrhythmia by buchang wenxin granule. *Henan Coll of Sci and Tech* (*Med Science*). 2009;02:119–120.
- 17. Jin HX, Huang T. 20 cases of clinical observation on Wenxin keli treating arrhythmia. *Chin Comm Doc.* 2007;(15):94.
- Xue JZ. 126 cases of clinical observation on Wenxin keli treating arrhythmia. *Chin J Integr Med Cardio*. 2008;3 (Suppl):103–104.
- Ren YH, Qiao HF. 46 cases of clinical observation on Wenxin keli treating arrhythmia. *Chin J Integr Med Cardio*. 2005;3(3):256–257.
- Wu XW, Yue L. 48 cases of clinical observation on Wenxin keli treating arrhythmia. *Chin J Integr Med Cardio*. 2004;2(8):487–488.
- Li ZY, Shen JX. 40 cases of clinical observation on Wenxin keli treating arrhythmia. *Zhejiang J Tradit Chin Med.* 2011;09:698.
- 22. Wang DQ. 40 cases of clinical observation on buchang Wenxin keli treating arrhythmia. *J Tradit Chin Med.* 2010;51(7):624.
- Pang YL. 56 cases of clinical observation on buchang Wenxin keli treating arrhythmia. *Yunnan J Tradit Chin Med Mater Medica*. 2010; 31(12):33.
- 24. Xu ZM, He YP, et al. 68 cases of clinical observation on buchang Wenxin keli treating arrhythmia. *J Chin Physician*. 2006;Suppl:562–563.
- Xia DM. Clinical efficacy observation on buchang Wenxin particles treating arrhythmias. *Modern Diagn Treat*. 2010;21(6):344–345.
- Sun XJ. 68 cases of clinical observation on buchang Wenxin keli treating ventricular contraction. *Liaoning J Tradit Chin Med.* 2009;09: 1530–1531.
- Wu Y. Clinical observation on Wenxin keli improving premature patient's symptoms and ECG. *Chin J Exp Med Formul.* 2009;8(15):92–93.
- Wang HQ. Clinical observation on the safety and efficacy of Wenxin keli treating ventricular contraction. *Clin Focus*. 2009;12:1082–1083.
- Guo YH. 112 cases of clinical observation on buchang Wenxin keli treating premature ventricular contractions. *Med Inf.* 2013;1(26):163.
- Jin RS. Clinical observation on buchang Wenxin keli treating ventricular contractions. *Chin Prac Med.* 2011;17(6):164–165.
- Zhang B, Liu F, Duan XX. Clinical observation on buchang Wenxin keli treating premature ventricular contractions. *Chin J Mod Drug Appl.* 2010;01:103–104.
- 32. Li ZM. 64 cases of clinical observation on buchang Wenxin keli treating premature ventricular contractions. *Chin Hth Nutrition*. 2012;12:243–244.
- Wang YJ. 60 cases of clinical observation on buchang Wenxin keli treating premature ventricular contractions. *Chin Prac Med.* 2011;16: 158–159.
- Yan GF. Clinical observation on treatment of 60 cases of premature ventricular vontraction with Wenxin granules. *Guid J Tradit Chin Med Pharm.* 2007;13(4):24–25.
- Yu YH, Li BH, et al. Clinical observation on buchang Wenxin keli treating angina. *Xinjiang J Tradit Chin Med Pharm*. 2004;04:14–15.
- Shu B, Li Y. Wenxin granules for treating 37 cases of angina pectoris of coronary atherosclerosis heart disease. *Chin Pharm*. 2013;22(16):102–103.
- Ye GF, Shi YF, et al. Clinical study of Wenxin granule on coronary heart disease with angina pectoris. *Jilin Med J.* 2008;29(5):362–363.
- Wei YQ, Zhang XW, et al. Clinical study of Wenxin keli treating unstable angina. Pract J Cardiac Cereb Pneum Vasc Dis. 2010;03:370–371.
- Yuan KY. Clinical analysis of Wenxin keli treating unstable angina. *Chin Prac Med.* 2011;22:141–142.
- 40. Wei YQ, Deng GY. Clinical study of Wenxin keli treating angina. *Liaoning J Tradit Chin Med.* 2010;08:1516.
- 41. Yuan JQ, Wei YY. Clinical study of Wenxin keli treating unstable angina. *Clin Med.* 2005;03:43–44.
- Yu T, Li DH, et al. 37 cases of clinical observation on Wenxin keli treating chronic heart failure. *Jilin J Tradit Chin Med.* 2006;11:9–10.

- Yang F, Dong ZH. Effect of Wenxin keli on chronic heart failure, plasma brain natriuretic peptide and cardiac function in patients. *Zhejiang J Tradit Chin Med*. 2009;01:72.
- Kong YH, Deng CJ, Cheng W, et al. Effect of Wenxin keli on cardiac function and plasma brain natrium peptide in patients with chronic heart failure. J Clin Cardiol. 2009;25(6):428–430.
- Xu KH. Clinical observation on Wenxin keli treating chronic heart failure. *Health for Everybody (Medical Guide)*. 2008;04:128.
- Hu JH. Effect of Wenxin granuleon chronic heart failure. *Chin Tradit* Patent Med. 2010;32(12):2035–2037.
- Yu XF. 37 cases of clinical observation on Wenxin keli treating heart failure. *Chin Prac Med.* 2008;05:95–96.
- Wang H. Clinical observation on Wenxin keli treating chronic congestive heart failure. *Liaoning J Tradit Chin Med.* 2012;11:2229–2230.
- Yang LL. Clinical observation of Wenxin granule in the treatment of children with viral myocarditis. *Modern Diagn Treat*. 2014;20: 4634–4635.
- Deng XC. Clinical analysis of Wenxin keli in treatment of children with viral myocarditis by assisting. *Yunnan J Tradit Chin Med Mater Medica*. 2012;07:86.

- Liu PX, Ren FX. 118 cases of clinical observation on Wenxin keli treating palpitations in patients with climacteric syndrome. *China Med.* 2008;8(3):493–494.
- Lei DF. 42 cases of clinical observation on Wenxin keli treating palpitations in patients with climacteric syndrome. *Shaanxi J Tradit Chin Med.* 2009;07:791–792.
- Li L, Miao F. Clinical observation on Wenxin keli treating climacteric syndrome. J Changzhi Med Coll. 2010;05:343–344.
- Hu XP. 35 cases of clinical observation on Wenxin keli treating climacteric syndrome. *Clin Res.* 2013;2:161.
- Qi P. 32 cases of clinical observation on Wenxin keli treating arrhythmia. *Yunnan J Tradit Chin Med Mater Medica*. 2010;08:49.
- Liu CL, Dong ZS. 32 cases of clinical observation on Wenxin keli treating tachyarrhythmias. *Chin J Integr Med Cardio*. 2005;3(1):68–69.
- 57. Ye F. 63 cases of clinical observation on Wenxin keli treating arrhythmia. *The Chin and Forgn Hth Abs.* 2013;10(10):204.

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