

The Efficacy of Guizhi Fuling Capsule or Kuntai Capsule Combined With Diane-35 and Metformin in Polycystic Ovary Syndrome: A Systematic Review and Meta-Analysis

Shuting Wei¹, Pu Ge², Haiming Hu³, Yudi Song⁴*

- 1.Translational Medicine Research Center, Medical Innovation Research Division and Fourth Medical Center of the Chinese PLA General Hospital, Beijing,100037, China
- 2. School of Traditional Chinese Medicine, Beijing University of Chinese Medicine, Beijing, 100029, China
- 3.Department of Pulmonary and Critical Care Medicine at The Seventh Medical Center, College of Pulmonary and Critical Care Medicine of The Eighth Medical Center, Chinese PLA General Hospital, Beijing, 100853, China
- 4. Shenzhen Maternity & Child Healthcare Hospital, Shenzhen, Guangdong, 518028, China
- *Corresponding author: Yudi Song, 17853140673@163.com

Copyright: 2025 Author(s). This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY-NC 4.0), permitting distribution and reproduction in any medium, provided the original author and source are credited, and explicitly prohibiting its use for commercial purposes.

Abstract: Background: Polycystic ovary syndrome (PCOS) is one of the most common reproductive endocrine metabolic diseases. Combined use of metformin and diane-35 has better curative effect in regulating serum hormone level (LH, FSH, T and E2) than using metformin alone. Traditional Chinese medicine (TCM) can also be used to treat PCOS. According to some studies, the combined use of metformin and diane-35 and TCM have achieved better curative effect than combining metformin and diane-35 in the treatment of patients with PCOS. Methods: Computerized searches of the science, Medline, VIP, Wan Fang and China HowNet (CNKI) databases were conducted to identify eligible randomized controlled trials (RCTs) from the data obtained up to March 1, 2022. The Cochrane Collaboration risk of bias tool was used to assess the risk of bias in individual RCTs, and R software(version 4.0.3) was used for data statistical analysis. Results: Nine RCTs involving 1035 patients were included. Comparing to D+M, significant reduce of LH(mean difference [MD]: -1.93, 95% confidence interval [CI]: -3.44, -0.42; Unit: U/L P < 0.01; I2=89%), T(MD: -1.44, 95%CI -2.59, -0.30; Unit: nmol/L P < 0.01; I2=98%) and significant increase of E2(MD: 31.43, 95%CI 24.54, 38.33;Unit:pmol/L P < 0.01; I2=96%) were shown in TCM+D+M. Comparing to D+M, TCM+D+M group has higher ovulation rate(RR 1.14 95%CI 1.07,1.22; P=0.42; I2=0%) and higher pregnancy rate(RR 1.29 95%CI 1.15,1.44; P=0.37; I2=7%). There is no significant difference between the two therapies in FSH changes (MD: -1.00, 95%CI -2.27, 0.28;Unit:U/L P < 0.01; I2=95%).Subgroup analysis showed that compared with the Guizhi Fuling capsule group, the Kuntai capsule group had more FSH reduction and E2 increase more. In other outcome indicators, the two subgroup did not show significant differences. Conclusion: Kuntai Capsule + Diane-35 + Metformin is better than Guizhi Fuling Capsule in reducing FSH, and it is also better in increasing E2. There was no significant difference between the two in LH and T hormones. There was no significant difference between Kuntai Capsules + Diane-35 + Metformin and Guizhi Fuling Capsules + Diane-35 + Metformin. As for the effect in lessen insulin resistance, Kuntai Capsule+Diane-35+Metformin was significantly better than Guizhi Fuling Capsules+Diane-35+Metformin.

Keywords: PCOS; Infertility; Guizhi Fuling Capsule; Kuntai Capsule; Diane-35; Metformin

Published: Aug 12, 2025

DOI: https://doi.org/10.62177/apjcmr.v1i3.516

1.Introduction

Polycystic ovary syndrome (PCOS) is one of the most common reproductive endocrine metabolic diseases, with a range of clinical manifestations such as oligo-anovulation, hyperandrogenism, metabolic abnormalities (insulin resistance and impaired fasting glucose levels), infertility and obesity^[1-6], affecting up to 20% of women of all ages and 5% to 10% of women of reproductive age^[5]. Although PCOS don't cause sudden deaths, it will greatly reduce the quality of life of women with the disease, for it can hardly be cured but to improve symptoms.

According to some studies, insulin resistance(IR) occurs in 70% of the patients with PCOS^[7-9]. International guidelines for the treatment of polycystic ovary system recommend metformin as the medicine for PCOS patients with IR. Some studies have shown that combined use of metformin and diane-35 has better curative effect in serum hormone level including Luteinizing hormone(LH), Follicle Stimulating Hormone (FSH), Testosterone (T), Estradiol(E2) than using metformin or diane-35 alone^[10-12].

In recent years, many studies have concentrated on the treatment of PCOS with traditional Chinese medicine(TCM), for instance, Kuntai capsule and Guizhi Fuling capsule have been confirmed effective for reducing the serum FSH and TSH level and ameliorating insulin resistance situation with little toxic side effects^[13-14]. Moreover, the RCT studies of Ming Luo^[15], Haiyan wang et,al^[16] and Xiu Li state that the combine use of Chinese and western medicine shows a better improvement of the PCOS patients in serum hormone level(LH,FSH,T or androgen) or conditions of ovulation and pregnancy than using the western medicine alone.

However, there are few systemic reviews and meta-analysis on the effect of combine use of TCM and western medicine on the treatment of PCOS. Qian-wen Ma et,al^[17]conducted a meta-analysis to conclude co-treatment with TCM and letrozole was more effective than letrozole monotherapy in the treatment of PCOS, but they didn't concentrate on certain kinds of TCM, so the conclusion can be difficult to put into clinical application, for as is well known, the prescription of some TCM can be variable and individual specific^[18].

Chinese patent medicine such as Kuntai capsule^[19] and Guizhi Fuling capsule^[14] have fixed formulations and recommended dosage, that is to say, the extensive clinical application and promotion of them can have realistic significance. However, up to date, no study has compared the effects of combining use of Kuntai capsule or Guizhi Fuling capsule and western medicine with using western medicine alone.

2. Objectives

The primary aim of this study was therefore to undertake a comprehensive systematic review and meta-analysis comparing the effect of Guizhi Fuling capsule/ Kuntai capsule + metformin + Diane-35 with metformin + Diane-35 in PCOS on a range of indexes including serum FSH, LT, E2, T levels and reproductive parameters including ovulation rate and pregnancy rate. We also conduct subgroup analyses to provide evidences for the choice of Guizhi Fuling capsule and Kuntai capsule as the TCM in the combine medication.

3. Materials and Methods

We followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses statement to conduct our systematic review. The full review protocol was registered with PROSPERO under registration number CRD 42021281360.

3.1 Study eligibility criteria

The PICO (population, intervention, comparison, outcome) framework was used to establish a priori selection criteria and included all RCTs comparing (i) Guizhi Fuling capsule + metformin + Diane-35 versus metformin + Diane-35, (ii) Kuntai capsule + metformin + Diane-35 versus metformin + Diane-35 for women of any age with PCOS.

Exclusion criteria:(i) the participants of the study were presence of other etiologies for hyperandrogenism or infertility, such as hypothyroidism, congenital adrenal hyperplasia and Cushing's syndrome, or concurrent medication use (e.g. OCPs, orlistat, clomiphene citrate, etc.) even if the same between groups;(ii)the study did not include the outcome measures we will

be studying; (iii)abstract data from conference and scarce literature; (iv) the quality evaluation of the study shows very poor quality.

Primary outcomes included metabolic parameters including serum hormone levels: Luteinizing hormone(LH), Follicle Stimulating Hormone (FSH), Testosterone (T), Estradiol(E2); and Secondary outcomes included reproductive parameters including ovulation rate and pregnancy rate.

3.2 Literature search methodology

English literature was searched in PubMed, web of science and Medline, and the literature in Chinese was searched in VIP, Wan Fang and China HowNet (CNKI) databases. There were no limits on year of publication. Language limit is Chinese and English.

The retrieval condition is as follows: "Chinese medicine", "Chinese herbal medicine" "traditional Chinese and western medicine", "Fuling Guizhi capsule/pill", "Guizhi Fuling capsule/pill", "Kuntai capsule/pill", "metformin", "Diane-35", "Ethinylestradiol cyproterone", and "Polysystic ovary syndrome", "PCOS" were searched in title, key words or abstract.

3.3 Study selection

In compliance with the inclusion criteria, we primarily screened the titles and abstracts, then retrieved the full texts of all potentially eligible studies. Two review authors independently examined and selected the eligible articles according to the inclusion criteria for the current review. Disagreement with respect to study eligibility was resolved via discussion with the third reviewer author.

3.4 Data extraction

Two review authors independently extracted the data from each eligible study. Any disagreements were resolved by discussion with a third review author. Data retrieved the basic characteristics of included study including sample size, mean age, interventions and duration time, as well as outcome measures. Authors were contacted for additional or missing information as required.

3.5 Risk of bias

The risk of bias of the included studies was assessed independently by two authors using the Cochrane Collaboration's tool for assessing risk of bias with respect to the following aspects: random sequence generation, allocation concealment, blinding of participants or personnel, blinding of outcome assessment, reporting bias, loss to follow up, other sources of bias. Any discrepancies were resolved via discussion with the third review author to reach a consensus.

3.6 Statistical analysis

Statistical processing was done using the statistical software package R (http://www.R-project.org/). The results were reported as mean difference (MD) with 95% CI for continuous outcomes of serum FSH, LT, E2, T levels and odds ratios (OR) with 95% confidence interval (95% CI) for dichotomous outcomes of ovulation rate and pregnancy rate. Subgroup analyses were performed based on the kind of TCM used in the integrated Chinese and Western medicine group. Publication bias was estimated by Egger's test^[20] in addition to funnel plot.

Heterogeneity between studies was examined using the I-square (I^2) index to quantify the degree of heterogeneity. If the test results present high heterogeneity ($I^2 \ge 50\%$), meta-analysis is performed using a random effect model; if the experimental results present low-moderate heterogeneity ($I^2 < 50\%$), meta-analysis is performed using a fixed effect model^[21].

4. Results

4.1 Study Selection and Characteristics

In the presence of heterogeneity, the two researchers checked the data entered and explored the variation by conducting sensitivity analysis. In the initial search, 370 relevant articles were identified, of which 130 articles were from CNKI, 92 articles from VIP, 139 articles from Wan Fang,2 articles were from pubmed, 2 articles from medline, and 5 articles from web of science. After the exclusion of 197 duplicate articles using EndNote X9 software, 173 articles underwent a title and abstract review. A total of 156 studies were excluded in this process, of which 12 articles are review studies,6 articles are animal research, the intervention methods of 138 articles did not meet the inclusion criteria. For the remaining 17 articles, the full text was reviewed and 8 of them were excluded for following reasons: the dosage form is decotion but not pill or

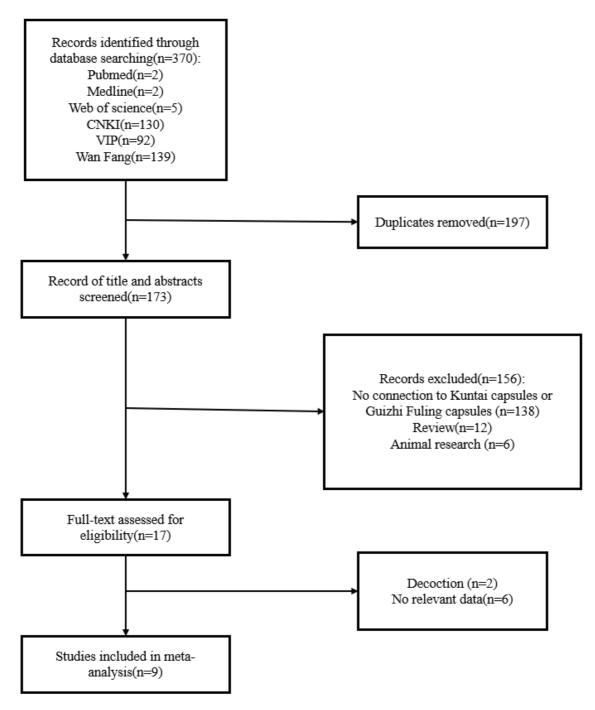
capsule(n= 2), which means that the other herbs may be added to the potion; article does not report the data required for this study (n= 6). The nine remaining studies fulfilled the eligibility criteria and were included in the meta-analysis $^{[13,15-16,22-27]}$. The literature review and identification process are shown in Figure 1. The meta-analysis evaluated the efficacy of two Chinese medicine preparations combined with Diane-35 and Metformin versus Diane-35 and Metformin for a total of 1035 patients with PCOS across nine clinical studies $^{[13,15-16,22-27]}$. Baseline clinical characteristics of these patients are summarized in Table 1.

Table 1 The characteristics from the eligible studies

	Sample size		Mean ag	ge(years)	Therapeutic r	nethod		
Study	Experi- mental	Con- trol	Experimen- tal	Control	Control Experimental		Outcome	Duration
Xiu Li, 2020	40	40	31.17±6.65	30.29±5.31	Diane-35+Met- formin+Cassia twig tuckahoe capsule	Di- ane-35+Met- formin	serum sex hormone levels(LH, FSH, E2, T); adverse reaction(trans- aminase,myocardial enzyme, creatinine, bone density); recrudesce	63 days
Liyun ZHANG, 2016	55	55	30. 8±6.9	29. 8±6.6	Diane-35+Met- formin+Cassia twig tuckahoe pill	Di- ane-35+Met- formin	serum sex hormone levels(LH, FSH, E2, T, Fins, FBG, TC, TG, HDL-C, LDL-C,); HOMA-IR; ISI; BMI; WHR; acne	Three men- strual cycles
Ying TIAN, 2017	54	53	29.8±4.5	29.3±4.1	Diane-35+Met- formin+Cassia twig tuckahoe capsule	Di- ane-35+Met- formin	serum sex hormone levels(LH, FSH, E2, T), HOMA-IR; HOMA-β, BMI; WHR; polytrichia; Ovarian volume; acne	Three menstrual cycles
Ming LUO, 2019	53	53	26.87±1.69	27.04±1.59	Diane-35+Met- formin+Kuntai capsule	Di- ane-35+Met- formin	serum sex hormone levels(LH, FSH, E2, T); HOMA-IR; HOMA-β	4 months
Haiyan WANG, 2020	80	80	26.50±1.73	26.19±1.54	Diane-35+Met- formin+Kuntai capsule	Di- ane-39+Met- formin	serum sex hormone levels(LH, FSH, E2, T); Ovarian volume; men- strual cycle	63 days
Qionglin LIN, 2016	49	49	26.53±1.46	26.61±1.47	Diane-35+Met- formin+Kuntai capsule	Di- ane-35+Met- formin	serum sex hormone levels(LH, FSH, E2, T); HOMA-IR; HOMA-β; Dizziness; headache; nausea; vomiting; skin irritation; breast tender	84 days
Hongmei Xv, 2020		52	26.96±1.49	26.80±1.45	Diane-35+Met- formin+Kuntai capsule	Di- ane-35+Met- formin	serum sex hormone levels(LH, FSH, E2, T); HOMA-IR; HOMA-β; polytrichia; acne;	Three men- strual cycles
Tong- shan HU, 2017	30	30	33.15±5.21	33.19±5.10	Diane-35+Met- formin+Kuntai capsule	Di- ane-35+Met- formin	HOMA-β; serum sex hormone(FSH); ovula- tion rate; pregnancy rate	100 days

	Sample	e size	Mean ag	ge(years)	Therapeutic r	nethod			
Study	Experi- mental	Con- trol	Experimen- tal	Control	Experimental	Control	Outcome	Duration	
Yan LIU, 2017	105	105	29.45±9.41	29.23±9.09	Diane-35+Met- formin+Kuntai capsule	Di- ane-35+Met- formin	serum sex hormone levels(LH, FSH, E2, T)	4 months	

Figure 1. Preferred Reporting Items for Systematic Reviews and Meta-Analyses flow diagram of the search process.

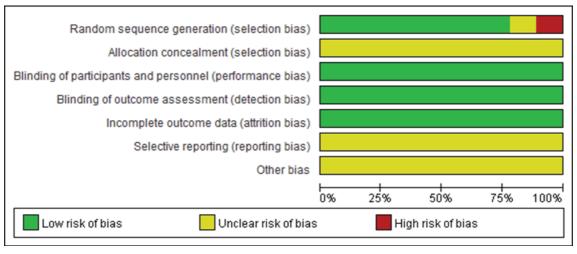


4.2 Quality Assessment

Trial quality was assessed using Cochrane risk of bias tool. In the nine studies, only one study were assessed as high risk of bias and no study were assessed as low risk of bias (Figure 2).

Blinding of participants and personnel (performance bias) Blinding of outcome assessment (detection bias) Random sequence generation (selection bias) Incomplete outcome data (attrition bias) Allocation concealment (selection bias) Selective reporting (reporting bias) Other bias ? Haiyan WANG 2020 Hongmei Xv 2020 ? ? ? ? Liyun ZHANG 2016 ? ? Ming LUO 2019 Qionglin LIN 2016 ? ? ? ? ? Tongshan HU 2017 ? ? ? Xiu Li 2020 Yan LIU 2017 ? ? Ying TIAN 2017

Figure 2. Assessment of risk of bias of included RCTs



4.3 Anthropometric parameters

4.3.1 Follicle Stimulating Hormone(FSH)

Seven studies reported the change in FSH involving 871 patients. There is no significant difference between the two therapies in FSH changes (MD:-1.00, 95%CI -2.27, 0.28;Unit:U/L; P < 0.01; $I^2=95\%$)(Figure 3).

Figure 3 Meta-analysis of TCM +D+M versus D+M for FSH(U/L). CI, confidence interval.

044	Experiment		M B///	Weight	Weight
Study	Total Mean S	D Total Mean SD	Mean Difference N	ID 95%-CI (common)	(random)
Xiu LI -2020	40 5.36 1.895	9 40 2.94 2.0797	2.	42 [1.55; 3.29] 5.8%	14.2%
Liyun ZHANG -2016	55 0.87 1.861	3 55 0.72 1.8570	0.	15 [-0.54; 0.84] 9.2%	14.5%
Ying TIAN-2017	54 -3.11 1.514	8 53 -1.88 3.4204	-1.	23 [-2.24; -0.22] 4.4%	13.9%
Ming LUO-2019	53 -9.98 3.812	2 53 -8.10 3.7995	-1.9	88 [-3.33; -0.43] 2.1%	12.7%
Haiyan WANG -2020	80 -2.66 1.296	9 80 -0.55 1.2271	-2.	11 [-2.50; -1.72] 28.9%	15.0%
Qionglin LIN-2016	49 -2.68 1.300	3 49 -0.51 1.2816	-2.	17 [-2.68; -1.66] 16.9%	14.8%
Yan LIU-2017	105 -2.60 1.438	9 105 -0.49 1.2808	-2.	11 [-2.48; -1.74] 32.6%	15.0%
Common effect model	436	435	-1.	61 [-1.82; -1.40] 100.0%	
Random effects model			-1.	00 [-2.27; 0.28]	100.0%
Heterogeneity: $I^2 = 95\%$, τ^2	2 = 2.7798, p < 0.01				
			-3 -2 -1 0 1 2 3		

4.3.2 Luteinizing hormone(LH)

Seven studies reported the change in LH involving 871 patients. Comparing to D+M, TCM +D+M significantly reduced LH(MD: -1.93, 95%CI -3.44, -0.42;Unit:U/L;P < 0.01; I^2 =89%)(Figure 4) .

Figure 4 Meta-analysis of TCM+D+M versus D+M for LH(U/L). CI, confidence interval.

Study	Total	Experi Mean	imental SD		(Mean	Control SD	Mean Difference	MD	95%-CI	Weight (common)	Weight (random)
Xiu LI -2020 Livun ZHANG -2016	40 55		4.1327 3.0925	40		4.5719 3.0780			[-7.14; -3.32]	3.4% 9.5%	12.6% 14.5%
Ying TIAN-2017	54	-11.94	3.2473 1.2987	53	-8.77	2.8324 1.2839		-3.17	[-2.72; -0.42] [-4.32; -2.02]	9.5% 9.5% 52.1%	14.5%
Ming LUO-2019 Haiyan WANG -2020	53 80	-7.54	3.9072		-9.38	3.8011]	1.84	[-2.60; -1.62] [0.65; 3.03]	8.8%	15.6% 14.4%
Qionglin LIN-2016 Yan LIU-2017	49 105		3.8422 3.8265			3.9368 3.9238			[-3.45; -0.37] [-2.80; -0.70]	5.3% 11.5%	13.6% 14.7%
Common effect model Random effects model Heterogeneity: $I^2 = 89\%$, τ^2	436 = 3.72	!52. p < !	0.01	435					[-2.22; -1.51] [-3.44; -0.42]	100.0%	100.0%
,-		, -					-6 -4 -2 0 2 4 6				

4.3.3 Estradiol(E2)

Seven studies reported the change in LH involving 871 patients. Comparing to D+M, TCM+D+M significantly increased E2(MD: 31.43, 95%CI 24.54, 38.33;Unit:pmol/L;P < 0.01; I^2 =96%)(Figure 5).

Figure 5 Meta-analysis of TCM +D+M versus D+M for E2(U/L). CI, confidence interval.

Study	Experimen Total Mean	al D Total Mea	Control n SD	Mean Difference	MD	95%-CI	Weight (common) (Weight random)
Xiu LI -2020 Liyun ZHANG -2016 Ying TIAN-2017 Ming LUO-2019 Haiyan WANG -2020 Qionglin LIN-2016 Yan LIU-2017	40 145.43 47.69 55 49.18 37.95 54 69.73 6.84 53 96.17 13.54 80 97.18 14.19 49 97.20 14.22 105 96.16 14.40	79 55 22.7 11 53 53.0 11 53 60.3 80 80 59.5 77 49 59.4	1 9.2684 8 9.2518	■ ■ = =	26.42 16.73 35.85 37.67 37.72	[1.17; 41.47] [12.44; 40.40] [13.92; 19.54] [31.45; 40.25] [33.96; 41.38] [32.97; 42.47] [33.78; 40.34]	0.6% 1.3% 32.0% 13.1% 18.3% 11.2% 23.5%	7.0% 10.2% 17.0% 16.3% 16.6% 16.1% 16.8%
Common effect model Random effects model Heterogeneity: I^2 = 96%, τ^2	436 = 70.9639, <i>p</i> < 0.01	435	! -4		30.35	[28.76; 31.94] [24.54; 38.33]	100.0%	 100.0%

4.3.4 Testosterone(T)

Six studies reported the change in T involving 765 patients. Comparing to D+M, TCM +D+M significantly reduced T(MD: -1.44, 95%CI -2.59, -0.30;Unit:nmol/L; P < 0.01; $I^2 = 98\%$)(Figure 6).

Figure 6 Meta-analysis of TCM +D+M versus D+M for T(nmol/L). CI, confidence interval.

		Exper	imental		(Control								Weight	Weight
Study	Total	Mean	SD	Total	Mean	SD		Mean	Differe	ence	M	D	95%-0	CI (common)	(random)
Xiu LI -2020	40	-2.04	0.8246	40	-0.64	0.7803		÷	: 1		-1.4	-1 0	1.75; -1.0	5] 16.9%	17.0%
Liyun ZHANG -2016	55	-0.33	0.4952	55	-0.32	0.5250			+				0.20; 0.1	•	17.1%
Ying TIAN-2017	54	-4.38	1.5464	53	-3.02	1.7346			łΤ				1.98; -0.7	•	16.4%
Haiyan WANG -2020	80	-0.41	1.7280	80	-0.25	1.7199			-		-0.1	6 [-	0.69; 0.3	7] 7.3%	16.6%
Qionglin LIN-2016	49	-5.03	1.7102	49	-3.20	1.7682					-1.8	3 [-	2.52; -1.1	4] 4.4%	16.2%
Yan LIU-2017	105	-5.22	1.7445	105	-1.27	1.8949	-		!		-3.9	5 [-	4.44; -3.4	6] 8.6%	16.7%
Common effect model	383			382					♦				0.89; -0.6	•	
Random effects model								-	=		1.4	4 [-2	2.59; -0.3	0]	100.0%
Heterogeneity: $I^2 = 98\%$, τ^2	= 1.98	361, <i>p</i> <	0.01				I	ı	ı	I	1				
							-4	-2	0	2	4				

4.4 Clinical outcomes

4.4.1 Ovulation rate

Six studies were included for the meta-analysis of ovulation rate involving 639 PCOS patients. Comparing to D+M, TCM +D+M group has higher ovulation rate(RR:1.14 95%CI 1.07,1.22; P=0.42; I²=0%)(Figure 7).

Figure 7 Meta-analysis of TCM+D+M versus D+M for ovulation rate(%). CI, confidence interval.

Study	Experime Events			ntrol Total	Risk Ratio	RR	95%-CI	Weight (common)	Weight (random)
Liyun ZHANG -2016	49	55	45	55	+ 11	1.09	[0.93; 1.27]	18.0%	17.1%
Ying TIAN -2017	49	54	39	53		1.23	[1.03; 1.48]	15.7%	12.4%
Haiyan WANG -2020	74	80	65	80		1.14	[1.01; 1.29]	26.0%	27.5%
Qionglin LIN -2016	46	49	39	49	<u> </u>	1.18	[1.01; 1.38]	15.6%	16.3%
Hongmei Xv -2020	48	52	45	52	 	1.07	[0.93; 1.22]	18.0%	23.3%
Tongshan HU -2017	26	30	17	30	<u> </u>	- 1.53	[1.09; 2.16]	6.8%	3.5%
Common effect model		320		319	🖐	1.16	[1.09; 1.24]	100.0%	
Random effects model Heterogeneity: $I^2 = 0\%$, τ^2		0.4	2		 	1.14	[1.07; 1.22]		100.0%
					0.5 1 2	-			

4.4.2 Pregnancy rate

Six studies were included for the meta-analysis of ovulation rate involving 639 PCOS patients. Comparing to D+M, TCM +D+M group has higher pregnancy rate(RR: 1.29 95%CI 1.15,1.44;P=0.37; I²=7%)(Figure 8).

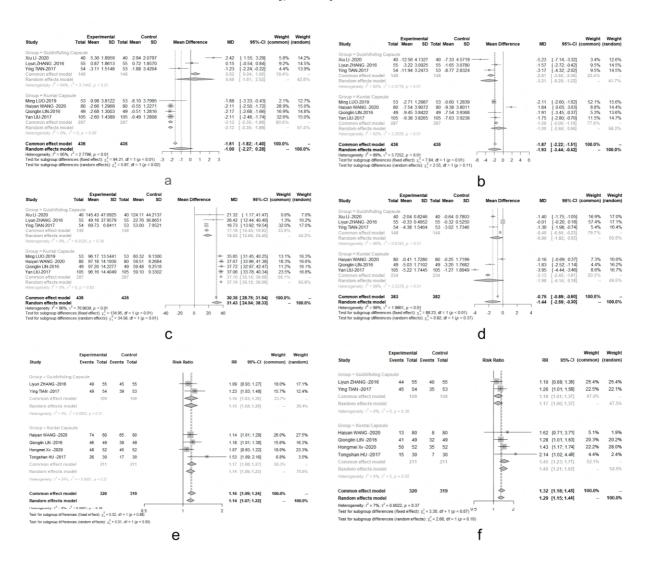
Figure 8 Meta-analysis of TCM+D+M versus D+M for pregnancy rate (%). CI, confidence interval.

Study	Experime Events			ntrol Total	Risk Ratio	RR	95%-CI	Weight (common)	Weight (random)
Liyun ZHANG -2016	44	55	40	55	1 2	1.10	[0.89; 1.36]	25.4%	25.4%
Ying TIAN -2017	45	54	35	53	 9 -	1.26	[1.01; 1.58]	22.5%	22.1%
Haiyan WANG -2020	13	80	8	80	- S =	1.62	[0.71; 3.71]	5.1%	1.9%
Qionglin LIN -2016	41	49	32	49	 	1.28	[1.01; 1.63]	20.3%	20.2%
Hongmei Xv -2020	50	52	35	52	=	1.43	[1.17; 1.74]	22.2%	28.0%
Tongshan HU -2017	15	30	7	30	-	- 2.14	[1.02; 4.49]	4.4%	2.4%
Common effect model		320		319	÷	1.32	[1.18; 1.48]	100.0%	
Random effects mode	•					1.29	[1.15; 1.44]		100.0%
Heterogeneity: $I^2 = 7\%$, τ^2	= 0.0022, p	0 = 0.37			1 1 1				
					0.5 1 2				

4.5 Subgroup analysis

To analyze the difference in efficacy of the two kinds of traditional Chinese medicine (Guizhifuling Capsule and Kuntai Capsule), we conducted subgroup analysis of all outcome indicators(Figure 9). Compared with the Guizhi Fuling capsule group, the Kuntai capsule group had more FSH reduction and E2 increase more, suggesting that Kuntai capsule may have better efficacy than Guizhi Fuling capsule. In other outcome indicators, the two groups did not show significant differences.

Figure 9 Subgroup analysis of G+D+M versus K+D+M for LSH(a), LH(b), E2(c), T(d), ovulation rate(e), and pregnancy rate(f). CI, confidence interval



4.6 Sensitivity Analysis

Because of the high heterogeneity, we evaluated the impact of every study on the pooled results to demonstrate stability and sensitivity (Figure 10-11). The results revealed that the outcomes of FSH and LH were reliable and stable.

Figure 10 Sensitivity analysis of FSH

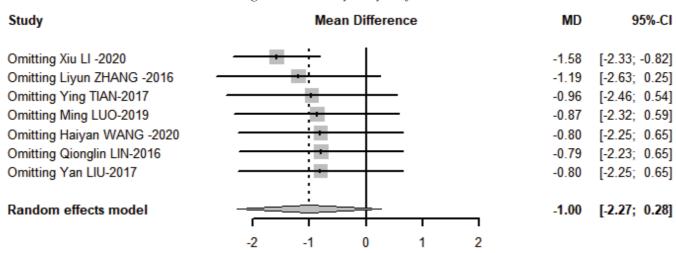


Figure 11 Sensitivity analysis of LH

Study	Mean Difference	MD	95%-CI
Omitting Xiu LI -2020	- ; = -	-1.46	[-2.81; -0.11]
Omitting Liyun ZHANG -2016		-2.00	[-3.79; -0.21]
Omitting Ying TIAN-2017	-	-1.73	[-3.45; 0.00]
Omitting Ming LUO-2019		-1.91	[-3.72; -0.09]
Omitting Haiyan WANG -2020	- :	-2.46	[-3.33; -1.59]
Omitting Qionglin LIN-2016		-1.94	[-3.73; -0.16]
Omitting Yan LIU-2017		-1.97	[-3.77; -0.17]
Random effects model		-1.93	[-3.44; -0.42]
	-3 -2 -1 0 1 2 3		

4.7 Publication Bias

The publication bias of the pooled LH(Egger's test: P = 0.9776), pooled pregnancy rate (Egger's test: P = 0.2543) were examined with funnel plot and Egger's regression tests (Figure 12-15). The results did not reveal evident publication bias in LH and pregnancy rate.

Figure 12 Funnel plot of LH

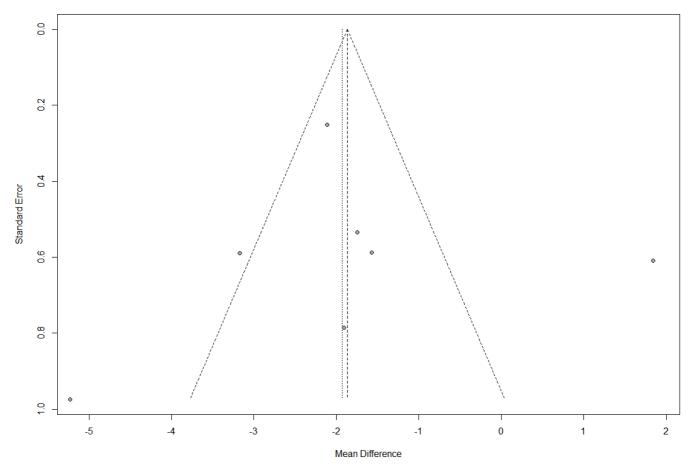


Figure 13 Egger's test of LH

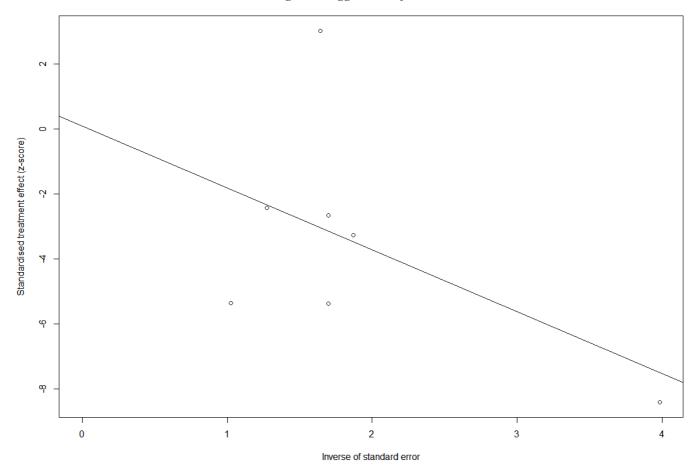


Figure 14 Funnel plot of pregancy rate

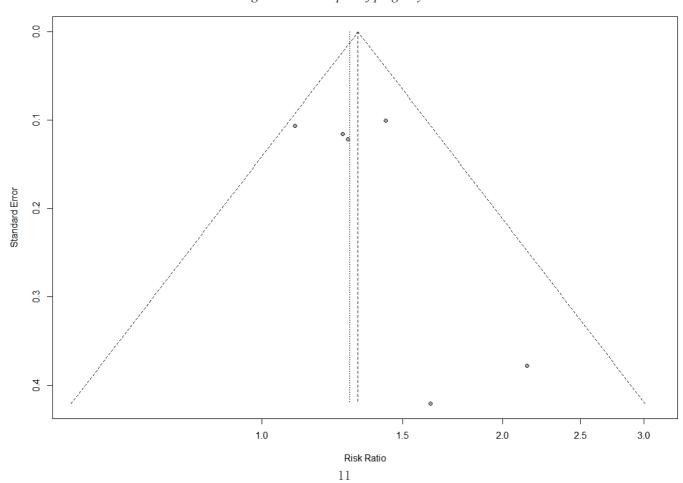
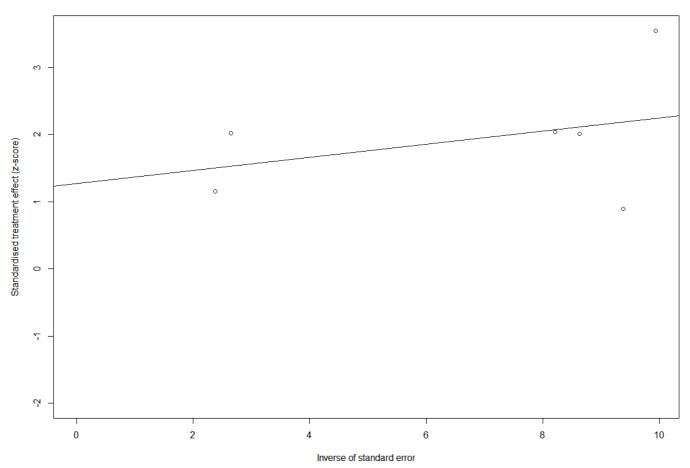


Figure 15 Egger's test of pregancy rate



5.Discussion

Polycystic ovary syndrome(PCOS) is the most common kind of endocrine disorder characterized by chronic anovulation, hyperandrogenism(HA) and insulin resistance(IR) and adversely affects the normal life of 6% to 20% of women in reproductive age globally^[28-29].

Some studies found that the excessive androgen lead to the inhibition of hypothalamic feedback, in other words, the elevated level of LH in serum cannot inhibit the further secretion of gonadotropin-releasing hormone(GnRH), which subsequently lead to the growing amount of $LH^{[30-31]}$. Excessive LH and low-dose FSH continuously stimulate the ovary, which contributing to the arrest of the growing of the dominant follicle.

However, the exact pathogenesis of PCOS still remains unclear, which result in the treatment strategy of this disease is dominated by symptomatic treatment, including weight control, life-style modification for the mild case and the regulation of menstrual cycle, anti-hyperandrogenemia (HA) therapy, management of insulin resistance(IR)and metabolic disorders for the severe one^[32-34]. One of the first line medication, Oral contraceptive pills (OCPs), have long been used to reduce HA level in serum and to regulate menstrual cycles in women with PCOS ^[35-36].OCPs not only possess the ability to increase the production of sex hormone-binding globulin (SHBG) to reduce serum free testosterone (T) hormone, the precursor of androgen, but also can reduce LH and androgen secretion by inhibiting hypothalamic gonadotropin^[37]. Diane-35, ethinyl estradiol (EE) plus cyproterone acetate (CPA), is the representative of OCPs and widely being used in China, Australia and Europe^[38].

As a biguanide used for the standard first-line treatment of type-2 diabetes mellitus, Metformin is applied in clinics to handle IR of patients with PCOS, it can enhance insulin sensitivity both in the liver and in the peripheral tissue through inhibiting hepatic glucose production and promoting uptake and utilization, respectively^[39-41]. A meta-analysis including 12 RCTs compared the effect of metformin +lifestyle change with placebo +lifestyle change, the result of which indicated that the former demonstrated more admirable effect in lowering BMI (MD: 20.73 kg/m2, 95% CI: 21.14, 20.32) and decreasing

subcutaneous adipose tissue (MD :292.49 cm², 95% CI: 2164.14, 220.84) and increasing the number of menstrual cycles (MD :1.06, 95% CI :0.30, 1.82) than the latter after 6 months intervention^[42]. Another meta-analysis focused on the Effect of metformin in overweight women also come to the similar conclusion that metformin reduces BMI, FSH, LH, low density lipoprotein (LDL), cholesterol and testosterone level in individuals with PCOS^[43].

An systematic review published in 2008 compared the efficacy of Diane-35 and metformin with Diane-35 alone, the result of which indicated that Diane-35 and metformin combination showed better result than Diane-35 monotherapy in lowering BMI(WMD 1.05; 95% CI 0.09, 2.01) in patients with PCOS^[38].

Some studies hypothesized that obesity might be more responsible than the elevated androgen level for the development of anovulation in PCOS and obesity induced metabolic abnormalities are closely related to insulin resistance(IR) [44-47]. When IR occurs in individuals, the function of insulin cannot be exerted which results in the synthesis and secretion of insulin from pancreatic cell being increased in a compensatory way^[48]. Hyperinsulinemia further deteriorates androgen-dependent anovulation via two main pathways^[48]. One is that insulin indirectly accelerates the androgen production through enhancing the effects of LH on ovarian theca cells^[49-50]. Another is that insulin inhibits the release of sex hormone-binding globulin (SHBG) from the liver, therefore the amount of bio-active androgen (unbound with SHBG) in blood circulation will increase^[51-52]. The excess androgen lead to menstrual irregularity, which in turn inhibit follicular maturation^[53]. Thus, obesity and IR will deteriorate hyperandrogenism in this feed-back circle, in addition to directly control the level of androgen, the active management of IR induced by obesity is of vital importance in the treatment of PCOS.

Western medicine treatment is mainly to adjust the menstrual cycle, improve ovulation, but none of them can permanently relieve the symptoms of patients, and even lead to ovarian overstimulation and low pregnancy rate^[54-56]. Moreover, metformin frequently induces gastrointestinal discomfort in patients, and in some cases even causes hypoglycemia, as dangerous as hyperglycemia, when uptake overdose or in fasting circumstances^[57-58].

Therefore, we intended to investigate whether the addition of Traditional Chinese Medicine(TCM) to metformin and Diane-35 would improve the clinical outcome of patients with PCOS.

TCM possess two major characteristic distinct from western medication, holistic therapy and multisystem regulation^[59].

In viewpoint of traditional Chinese medicine (TCM), the disorders of the kidneys, liver and spleen might result in the occurrence of PCOS, the deficiency in kidney in particular, since kidney is believed to govern reproductive function of humans^[60-62]. When the function of the liver is impaired, the blood circulation and the menstrual cycle of an individual will be adversely affected, while the condition of spleen will influence body type, obesity might occur if it doesn't perform well^[63].

Guizhi fuling capsule (GF) is one of the most frequently prescribed Chinese herbal formulas for treating uterine fibroids and widely accepted for the management of gynaecological conditions^[64]. The ingredients of guizhi fuling capsule consist of the following 5 herbs: Cinnamomum cassia (L.) J. Presl (Guizhi), Poria cocos (Schw.) Wolf (Fuling), Paeonia lactiflora Pall. (Chishao), Juglans regia L. (Taoren), and Paeonia suffruticosa Andr. (Mudanpi)^[65].

Some studies indicated that GF might ameliorate PCOS-IR through regulating related target proteins and pathways in hormone generation and secretion [66-68]. The function of GF is not limited to alleviating IR resistance, reducing fasting blood glucose together with insulin and correcting abnormal lipid metabolism, can also shorten the menstrual cycle, restore ovulation, and improve the pregnancy rate [69-71].

Another kind of the traditional Chinese medicine, Kuntai capsule (KT), consists of prepared Dihuang (Rehmannia glutinosa (Gaertn.) DC.), Huanglian (Rhizoma Coptidis), Baishao (Radix Paeoniae Alba), Huangqin (Radix Scutellariae Baicalensis), Ejiao (Colla Corii Asini), and Fuling (Poria)^[72]. The result of a meta-analysis demonstrated that Kuntai capsule has better performance in regulating serum sex hormone levels including LH, LH/FSH and T than placebo in patients with PCOS^[72].

In the mechanism analysis of the modern pharmacology, the component dang gui and baishao may reduce the release of insulin and androgen through phosphatidylinositol 3-hydroxy kinase(PI3K)/protein kinase B (AKT)/glucose transporter 4 (GLUT4) signal pathway and oxidative stress^[73-75].

Studies have shown that traditional Chinese medicine can inhibit the apoptosis and excessive autophagy of ovarian granulosa cells, reduce follicular atresia, improve ovarian reserve function, regulate serum endocrine hormone levels, restore the

number of ovarian granulosa cells, so as to maintain ovarian function and promote follicle development^[76-77].

The pooled results indicated that the efficacy of TCM+ diane-35+ metformin was significantly superior to that of diane-35+ metformin in lowering LH level (MD: -1.93, 95%CI: -3.44, -0.42;Unit:U/L; P < 0.01; I²=89%) and T level (MD: -1.44, 95%CI:-2.59,-0.30;Unit:nmol/L; P < 0.01; I²=98%)and increasing Estradiol(E2)(MD: 31.43, 95%CI 24.5,38.3;Unit:pmol/L; P < 0.01; I²=96%),Ovulation rate(RR 1.14 95%CI: 1.1,1.2; P =0.42; I²=0%)and Pregnancy rate(RR 1.29 95%CI 1.2,1.4; P =0.37; I²=7%).

A similar result can be find in a systematic review, which concludes that traditional Chinese medicine (TCM) has the competence to significantly increase the pregnancy rate and the ovulation rate in participants struggling with infertility due to anovulation^[78].

In attempt to find out whether there was some difference towards the type of TCM ,we conducted subgroup analysis stratified studies in according to the type of TCM.KT achieved better performance in increasing Estradiol (E2) (pooled MD: 31.4, 95%CI:24.5,38.3, Unit: pmol/L, P<0.01; $I^2=95.6\%$) and lowering FSH (pooled MD: -1.00,95%CI: -2.27,0.28;Unit: pmol/L, P=0.02; $I^2=95\%$) in comparison with GF.

Only one study reported the incidence of drug related adverse events^[23], as mentioned by which ,there were no severe adverse events in both groups and occasionally some patients developed slight increase in transaminase, myocardial enzyme and creatinine, which return to normal after two weeks. As a consequence, the pooled analysis is only conducted to the efficacy, while toxicity hasn't undergone pooled analysis.

6.Limitation

Limitation of our studies include: first, the included studies have not reported the diagnostic criteria and important outcome indicator like the live birth rate and adverse events, which result in the limited information was obtained in this meta-analysis; Second, although we conducted comprehensive search, included studies were all carried out in China, the result of the current meta-analysis might cannot be applied to the other part of the world^[79]; last, we suppose there might exist methodological flows since neither of them reported the calculation of sample size in the initiation and the drop-out rate in the end.

7. Conclusion

The results of our systematic review and meta-analysis validated the efficacy of TCM including GF and KT in the treatment of patients with PCOS. TCM containing therapy might be a potential option to enhance the efficacy of metformin and Diane-35. However, since the number of eligible studies is limited and most of them are relatively moderate quality, the results should be interpreted with caution. To provide stronger evidence for the application of GF or HYKT-containing TCM in the treatment of PCOS, more high quality randomized controlled trials are needed.

Funding

no

Conflict of Interests

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

Reference

- [1] Oerlemans, S., Mols, F., Nijziel, M. R., Lybeert, M., & van de Poll-Franse, L. V. (2011). The impact of treatment, socio-demographic and clinical characteristics on health-related quality of life among Hodgkin's and non-Hodgkin's lymphoma survivors: a systematic review. Annals of Hematology, 90, 993-1004.
- [2] Wang, A., Mo, T., Li, Q., Shen, C., & Liu, M. (2019). The effectiveness of metformin, oral contraceptives, and lifestyle modification in improving the metabolism of overweight women with polycystic ovary syndrome: a network meta-analysis. Endocrine, 64, 220-232.
- [3] Bozdag, G., Mumusoglu, S., Zengin, D., Karabulut, E., & Yildiz, B. O. (2016). The prevalence and phenotypic features of polycystic ovary syndrome: a systematic review and meta-analysis. Human reproduction, 31(12), 2841-2855.

- [4] Amiri, M., Kabir, A., Nahidi, F., Shekofteh, M., & Ramezani Tehrani, F. (2018). Effects of combined oral contraceptives on the clinical and biochemical parameters of hyperandrogenism in patients with polycystic ovary syndrome: a systematic review and meta-analysis. The European Journal of Contraception & Reproductive Health Care, 23(1), 64-77.
- [5] Carmina, E., & Lobo, R. A. (1999). Polycystic ovary syndrome (PCOS): arguably the most common endocrinopathy is associated with significant morbidity in women. The journal of clinical endocrinology & metabolism, 84(6), 1897-1899.
- [6] Norman, R. J., Dewailly, D., Legro, R. S., & Hickey, T. E. (2007). Polycystic ovary syndrome. The Lancet, 370(9588), 685-697.
- [7] Hackbart, K. S., Cunha, P. M., Meyer, R. K., & Wiltbank, M. C. (2013). Effect of glucocorticoid-induced insulin resistance on follicle development and ovulation. Biology of reproduction, 88(6), 153-1.
- [8] Farrell, K., & Antoni, M. H. (2010). Insulin resistance, obesity, inflammation, and depression in polycystic ovary syndrome: biobehavioral mechanisms and interventions. Fertility and sterility, 94(5), 1565-1574.
- [9] Legro, R. S., Castracane, V. D., & Kauffman, R. P. (2004). Detecting insulin resistance in polycystic ovary syndrome: purposes and pitfalls. Obstetrical & gynecological survey, 59(2), 141-154.
- [10] Han, Y., Li, Y., & He, B. (2019). GLP-1 receptor agonists versus metformin in PCOS: a systematic review and meta-analysis. Reproductive biomedicine online, 39(2), 332-342.
- [11] Feng, W., Jia, Y. Y., Zhang, D. Y., & Shi, H. R. (2016). Management of polycystic ovarian syndrome with Diane-35 or Diane-35 plus metformin. Gynecological Endocrinology, 32(2), 147-150.
- [12] Li, X., Guo, Y. R., Lin, J. F., Feng, Y., Billig, H., & Shao, R. (2014). Combination of Diane-35 and metformin to treat early endometrial carcinoma in PCOS women with insulin resistance. Journal of Cancer, 5(3), 173.
- [13] Hu, T. S. (2017). Clinical efficacy analysis of Kuntai capsule combined with ethinylestradiol cyproterone tablets and metformin hydrochloride tablets in the treatment of polycystic ovary syndrome. Electronic Journal of Clinical Medical Literature, 4(97), 19180–19181.
- [14] Liu, M., Zhu, H., Hu, X., Zhu, Y., & Chen, H. (2020). Efficacy of different forms of Guizhi Fuling Wan on reproduction and metabolism in women with polycystic ovary syndrome: a protocol for a meta-analysis of randomized controlled trials. Medicine, 99(44), e22954.
- [15] Luo, M.(2019). Effects of Kuntai Capsule adjuvant therapy on serum hormones and insulin resistance in patients with polycystic ovary syndrome. Capital Food and Medicine, 26(7), 64.
- [16] Wang, H. Y., Bi, X. L., Cao, J., & Feng, Q. (2020). Effects of Kuntai capsule combined with ethinylestradiol cyproterone tablets and metformin in the treatment of polycystic ovary syndrome and its impact on sex hormones. Medical Information, 33(20), 129–131.
- [17] Ma, Q. W., & Tan, Y. (2017). Effectiveness of co-treatment with traditional Chinese medicine and letrozole for polycystic ovary syndrome: a meta-analysis. Journal of Integrative Medicine, 15(2), 95-101.
- [18] Li, X., Wu, Z., Du, X., Wu, Y., Xie, X., & Shi, L. (2021). Interventions for preventing cardiotoxicity in breast cancer patients receiving trastuzumab: a systemic review and Bayesian network meta-analysis. Frontiers in Pharmacology, 12, 718086.
- [19] Liu, Y., Ma, H. X., & Yu, C. Y. (2019). A systematic review of Kuntai capsule in the treatment of infertility caused by polycystic ovary syndrome. Guangxi Journal of Traditional Chinese Medicine, 42(3), 59–65.
- [20] Egger, M., Smith, G. D., Schneider, M., & Minder, C. (1997). Bias in meta-analysis detected by a simple, graphical test. BMJ, 315(7109), 629-634.
- [21] Higgins, J. P. (2008). Cochrane handbook for systematic reviews of interventions. Cochrane Collaboration and John Wiley & Sons Ltd.
- [22] Xu, H. M. (2020). Clinical study of Kuntai capsule combined with ethinylestradiol cyproterone tablets and metformin in the treatment of polycystic ovary syndrome. Northern Pharmacy, 17(5), 36–37.
- [23] Li, X.(2020). Clinical efficacy observation of Guizhi Fuling capsule combined with ethinylestradiol cyproterone tablets and metformin in the treatment of polycystic ovary syndrome. China Practical Medicine, 15(7), 160–162. https://doi.

- org/10.14163/j.cnki.11-5547/r.2020.07.070
- [24] Lin, Q. L. (2016). Clinical study of Kuntai capsule combined with ethinylestradiol cyproterone tablets and metformin in the treatment of polycystic ovary syndrome. Drugs & Clinic, 31(3), 338–341.
- [25] Liu, Y.(2017). Application value of Kuntai capsule in the treatment of obesity-related polycystic ovary syndrome. International Journal of Women's Health Research, (8), 90, 103.
- [26] Tian, Y., & Gao, X. L.(2017). Effects of Guizhi Fuling capsule combined with western medicine on polycystic ovary syndrome: Impacts on endocrine metabolism and ovulation. Shaanxi Journal of Traditional Chinese Medicine, 38(4), 444–445.
- [27] Zhang, L. Y., & Yin, W. Q. (2016). Effects of Guizhi Fuling pill adjuvant therapy on patients with polycystic ovary syndrome and insulin resistance. Journal of Chinese Medicinal Materials, 39(7), 1661–1663. https://doi.org/10.13863/j.issn1001-4454.2016.07.051
- [28] Yildiz, B. O., Bozdag, G., Yapici, Z., Esinler, I., & Yarali, H. (2012). Prevalence, phenotype and cardiometabolic risk of polycystic ovary syndrome under different diagnostic criteria. Human reproduction, 27(10), 3067-3073.
- [29] Mohammad, M. B., & Seghinsara, A. M. (2017). Polycystic ovary syndrome (PCOS), diagnostic criteria, and AMH. Asian Pacific journal of cancer prevention: APJCP, 18(1), 17.
- [30] Blank, S. K., McCartney, C. R., & Marshall, J. C. (2006). The origins and sequelae of abnormal neuroendocrine function in polycystic ovary syndrome. Human reproduction update, 12(4), 351-361.
- [31] McCartney, C. R., & Marshall, J. C. (2016). Polycystic ovary syndrome. New England Journal of Medicine, 375(1), 54-64.
- [32] Song, S. Y., Yang, J. B., Song, M. S., Oh, H. Y., Lee, G. W., Lee, M., ... & Yoo, H. J. (2019). Effect of pretreatment with combined oral contraceptives on outcomes of assisted reproductive technology for women with polycystic ovary syndrome: a meta-analysis. Archives of Gynecology and Obstetrics, 300, 737-750.
- [33] Naderpoor, N., Shorakae, S., de Courten, B., Misso, M. L., Moran, L. J., & Teede, H. J. (2015). Metformin and lifestyle modification in polycystic ovary syndrome: systematic review and meta-analysis. Human reproduction update, 21(5), 560-574.
- [34] Abu Hashim, H., Foda, O., & El Rakhawy, M. (2018). Unilateral or bilateral laparoscopic ovarian drilling in polycystic ovary syndrome: a meta-analysis of randomized trials. Archives of gynecology and obstetrics, 297, 859-870.
- [35] Helvaci, N., & Yildiz, B. O. (2014). Oral contraceptives in polycystic ovary syndrome. Minerva endocrinologica, 39(3), 175-187.
- [36] Rojanasakul, A., Sirimongkolkasem, R., Piromsawasdi, S., Sumavong, V., Chailurkit, L. O., & Chaturachinda, K. (1987). Effects of combined ethinylestradiol, and desogestrel on hormone profiles and sex hormone binding globulin in women with polycystic ovarian disease. Contraception, 36(6), 633-640.
- [37] Carlström, K., Karlsson, R., & Schoultz, B. V. (2002). Diurnal rhythm and effects of oral contraceptives on serum dehydroepiandrosterone sulfate (DHEAS) are related to alterations in serum albumin rather than to changes in adrenocortical steroid secretion. Scandinavian Journal of Clinical and Laboratory Investigation, 62(5), 361-368.
- [38] Jing, Z., Liang-Zhi, X., Tai-Xiang, W., Ying, T., & Yu-Jian, J. (2008). The effects of Diane-35 and metformin in treatment of polycystic ovary syndrome: an updated systematic review. Gynecological Endocrinology, 24(10), 590-600.
- [39] Barbieri, R. L., Makris, A., Randall, R. W., Daniels, G., Kistner, R. W., & RYAN, K. J. (1986). Insulin stimulates androgen accumulation in incubations of ovarian stroma obtained from women with hyperandrogenism. The Journal of Clinical Endocrinology & Metabolism, 62(5), 904-910.
- [40] Dunn, C. J., & Peters, D. H. (1995). Metformin: a review of its pharmacological properties and therapeutic use in non—insulin-dependent diabetes mellitus. Drugs, 49, 721-749.
- [41] Nardo, L. G., & Rai, R. (2001). Metformin therapy in the management of polycystic ovary syndrome: endocrine, metabolic and reproductive effects. Gynecological endocrinology, 15(5), 373-380.
- [42] Domecq, J. P., Prutsky, G., Mullan, R. J., Hazem, A., Sundaresh, V., Elamin, M. B., ... & Murad, M. H. (2013). Lifestyle

- modification programs in polycystic ovary syndrome: systematic review and meta-analysis. The Journal of Clinical Endocrinology & Metabolism, 98(12), 4655-4663.
- [43] Guan, Y., Wang, D., Bu, H., Zhao, T., & Wang, H. (2020). The effect of metformin on polycystic ovary syndrome in overweight women: a systematic review and meta-analysis of randomized controlled trials. International journal of endocrinology, 2020(1), 5150684.
- [44] Rosenfield, R. L., & Ehrmann, D. A. (2016). The pathogenesis of polycystic ovary syndrome (PCOS): the hypothesis of PCOS as functional ovarian hyperandrogenism revisited. Endocrine reviews, 37(5), 467-520.
- [45] Alvarez-Blasco, F., Botella-Carretero, J. I., San Millán, J. L., & Escobar-Morreale, H. F. (2006). Prevalence and characteristics of the polycystic ovary syndrome in overweight and obese women. Archives of internal medicine, 166(19), 2081-2086.
- [46] Dunaif, A. (1997). Insulin resistance and the polycystic ovary syndrome: mechanism and implications for pathogenesis. Endocrine reviews, 18(6), 774-800.
- [47] Rojas, J., Chávez, M., Olivar, L., Rojas, M., Morillo, J., Mejías, J., Calvo, M., & Bermúdez, V. (2014). Polycystic ovary syndrome, insulin resistance, and obesity: navigating the pathophysiologic labyrinth. International journal of reproductive medicine, 2014, 719050. https://doi.org/10.1155/2014/719050
- [48] Sanchez-Garrido, M. A., & Tena-Sempere, M. (2020). Metabolic dysfunction in polycystic ovary syndrome: Pathogenic role of androgen excess and potential therapeutic strategies. Molecular metabolism, 35, 100937.
- [49] Baillargeon, J. P., & Carpentier, A. (2007). Role of insulin in the hyperandrogenemia of lean women with polycystic ovary syndrome and normal insulin sensitivity. Fertility and sterility, 88(4), 886-893.
- [50] Baillargeon, J. P., & Nestler, J. E. (2006). Polycystic ovary syndrome: a syndrome of ovarian hypersensitivity to insulin?. The Journal of Clinical Endocrinology & Metabolism, 91(1), 22-24.
- [51] Nestler, J. E., Powers, L. P., Matt, D. W., Steingold, K. A., Plymate, S. R., Rittmaster, R. S., ... & BLACKARD, W. G. (1991). A direct effect of hyperinsulinemia on serum sex hormone-binding globulin levels in obese women with the polycystic ovary syndrome. The Journal of clinical endocrinology & metabolism, 72(1), 83-89.
- [52] PLYMATE, S. R., MATEJ, L. A., JONES, R. E., & FRIEDL, K. E. (1988). Inhibition of sex hormone-binding globulin production in the human hepatoma (Hep G2) cell line by insulin and prolactin. The Journal of Clinical Endocrinology & Metabolism, 67(3), 460-464.
- [53] Nestler, J. E., Jakubowicz, D. J., Falcon de Vargas, A., Brik, C., Quintero, N., & Medina, F. (1998). Insulin stimulates testosterone biosynthesis by human thecal cells from women with polycystic ovary syndrome by activating its own receptor and using inositolglycan mediators as the signal transduction system. The Journal of Clinical Endocrinology & Metabolism, 83(6), 2001-2005.
- [54] Wang, R., Li, W., Bordewijk, E. M., Legro, R. S., Zhang, H., Wu, X., ... & International Ovulation Induction IPDMA Collaboration. (2019). First-line ovulation induction for polycystic ovary syndrome: an individual participant data meta-analysis. Human reproduction update, 25(6), 717-732.
- [55] Bordewijk, E. M., Ng, K. Y. B., Rakic, L., Mol, B. W. J., Brown, J., Crawford, T. J., & van Wely, M. (2020). Laparoscopic ovarian drilling for ovulation induction in women with anovulatory polycystic ovary syndrome. Cochrane Database of Systematic Reviews, (2).
- [56] Di Pietro, M., Velazquez, C., Matzkin, M. E., Frungieri, M. B., Peña, M. G., de Zúñiga, I., ... & Abramovich, D. (2020).
 Metformin has a direct effect on ovarian cells that is dependent on organic cation transporters. Molecular and Cellular Endocrinology, 499, 110591.
- [57] Palomba, S., Falbo, A., Zullo, F., & Orio Jr, F. (2009). Evidence-based and potential benefits of metformin in the polycystic ovary syndrome: a comprehensive review. Endocrine reviews, 30(1), 1-50.
- [58] Eisenhardt, S., Schwarzmann, N., Henschel, V., Germeyer, A., Von Wolff, M., Hamann, A., & Strowitzki, T. (2006). Early effects of metformin in women with polycystic ovary syndrome: a prospective randomized, double-blind, placebo-controlled trial. The Journal of Clinical Endocrinology & Metabolism, 91(3), 946-952.

- [59] Zhang, J., Li, T., Zhou, L., Tang, L., Xu, L., Wu, T., & Lim, D. C. (2010). Chinese herbal medicine for subfertile women with polycystic ovarian syndrome. Cochrane Database of Systematic Reviews, (9).
- [60] Ni, H. Y., & Gong, J.(2007). Research progress in traditional Chinese medicine treatment of polycystic ovary syndrome. Liaoning Journal of Traditional Chinese Medicine, (1), 123–124. https://doi.org/10.13192/j.ljtcm.2007.01.127.nihy.083
- [61] Wang, B. Q., & Lin, M.(2008). Research progress in traditional Chinese medicine for polycystic ovary syndrome. Shandong Journal of Traditional Chinese Medicine, 27(2), 138–140. https://doi.org/10.16295/j.cnki.0257-358X.2008.02.023
- [62] Huang, C. J., & Zhao, W.(2019). Professor Zhao Wei's clinical experience in treating infertility caused by polycystic ovary syndrome. Journal of Zhejiang Chinese Medical University, 43(4), 346–349. https://doi.org/10.16466/j.issn1005-5509.2019.04.013
- [63] Hou, F., & Shi, Y. Q.(2012). Research progress on etiology of polycystic ovary syndrome in traditional Chinese and Western medicine. Journal of Changchun University of Chinese Medicine, 28(5), 932–934. https://doi.org/10.13463/j.cnki.cczyy.2012.05.089
- [64] Yen, H. R., Chen, Y. Y., Huang, T. P., Chang, T. T., Tsao, J. Y., Chen, B. C., & Sun, M. F. (2015). Prescription patterns of Chinese herbal products for patients with uterine fibroid in Taiwan region: a nationwide population-based study. Journal of ethnopharmacology, 171, 223-230.
- [65] Li, M., Hung, A., & Yang, A. W. H. (2019). Guizhi Fuling Wan for uterine fibroids: A systematic review of in vivo studies. Journal of Ethnopharmacology, 245, 112177.
- [66] Shao, X. M.(2013). Clinical study of berberine combined with Guizhi Fuling Pill in patients with polycystic ovary syndrome and insulin resistance. Chinese Journal of Clinical Research, 26(8), 803–805.
- [67] Cao, L. H., Zhang, H. Q., & Zhao, Q.(2017). Effects of Guizhi Fuling Pill combined with metformin on inflammatory response and oxidative stress related to insulin resistance in PCOS patients. Journal of Hainan Medical University, 23(23), 3211–3214, 3218. https://doi.org/10.13210/j.cnki.jhmu.20171127.002
- [68] Yu, Y., Zhang, G., Han, T., & Huang, H. L.(2019). Mechanism of action of Guizhi Fuling Pill in treating polycystic ovary syndrome based on network pharmacology and bioinformatics analysis. Chinese Journal of Pharmacology and Toxicology, 33(10), 832–833.
- [69] Zhao, Q. S., Tan, X. F., & Wang, N. S.(2012). Effects of Guizhi Fuling Pill on insulin resistance and adiponectin in rats with polycystic ovary syndrome. Journal of New Chinese Medicine, 44(1), 116–117. https://doi.org/10.13457/j.cnki.jncm.2012.01.058
- [70] Tao, L., Tuo, A. X., & Liu, M. Y.(2013). Therapeutic effects of Guizhi Fuling Pill combined with berberine in polycystic ovary syndrome patients with insulin resistance. Chinese Journal of Experimental Traditional Medical Formulae, 19(15), 320–323.
- [71] Wu, J. J., Deng, H. T., & Liu, R.(2020). Clinical research progress of Guizhi Fuling Pill in treating infertility. Journal of Guangzhou University of Chinese Medicine, 37(3), 586–590. https://doi.org/10.13359/j.cnki.gzxbtcm.2020.03.038
- [72] Liang, R., Liu, Z., Li, P., Fan, P., Xu, L., Sun, X., ... & Zhang, M. (2019). Kuntai capsules improve glucolipid metabolism in patients with polycystic ovary syndrome: a randomized, double-blind, placebo-controlled trial. Medicine, 98(39), e16788.
- [73] Wang, X. C., & Zhang, Q. P.(2012). Research progress on treating polycystic ovary syndrome with the method of tonifying kidney, clearing liver and activating blood. Zhejiang Journal of Traditional Chinese Medicine, 47(3), 232–233.
- [74] Meng, X. Y., Guo, S. M., & Yang, L. X.(2017). Research progress on the mechanism of plant polysaccharides from traditional Chinese medicine for insulin resistance in type 2 diabetes. Chinese Journal of Experimental Traditional Medical Formulae, 23(8), 220–225. https://doi.org/10.13422/j.cnki.syfjx.2017080220
- [75] Peng, X. J., Xu, H. Y., Chen, Y. B., Yang, C. H., Xu, G., Liu, Y. H., & Yang, X. J.(2019). Network pharmacology analysis of the mechanism of Danggui Liuhuang Tang in treating diabetes. Traditional Chinese Drug Research and Clinical Pharmacology, 30(8), 952–958. https://doi.org/10.19378/j.issn.1003-9783.2019.08.010
- [76] Bai, J., Gao, R. R., & Wu, K. M.(2017). Research progress and review on the correlation between autophagy and

- delaying ovarian aging with traditional Chinese medicine. Chinese Archives of Traditional Chinese Medicine, 35(12), 3117–3120. https://doi.org/10.13193/j.issn.1673-7717.2017.12.035
- [77] Yang, S. W., Guo, J. S., Wang, X. Q., & Liu, D. D. (2015). Experimental study on Bushen Liaogeng extract delaying ovarian function decline in aged mice. Traditional Chinese Drug Research and Clinical Pharmacology, 26(5), 613–618.
- [78] Morley, L. C., Tang, T., Yasmin, E., Norman, R. J., & Balen, A. H. (2017). Insulin-sensitising drugs (metformin, rosiglitazone, pioglitazone, D-chiro-inositol) for women with polycystic ovary syndrome, oligo amenorrhoea and subfertility. Cochrane Database of Systematic Reviews, (11).
- [79] Gao, J., Zhang, W. J., & Yang, F. (2022). A Review of Traditional Chinese Medicine for the Treatment of Depression. Psychosom Med Res, 4(1), 6.