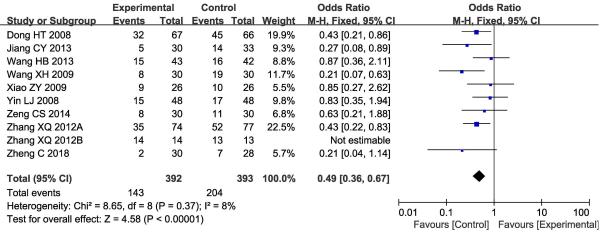


**Supplementary Figure 1.** Risk of bias summary: review of authors' judgments about each risk of bias item for included studies. **Note:** Each color represents a different level of bias: red for high-risk, green for low-risk, and yellow for unclear-risk of bias.





 $\mathbf{B}$ 

	Experim	ental	Contr	ol		Odds Ratio	Odds Ratio
Study or Subgroup	<b>Events</b>	Total	<b>Events</b>	Total	Weight	M-H, Fixed, 95% C	I M-H, Fixed, 95% CI
Jiang CY 2013	6	30	13	33	21.4%	0.38 [0.12, 1.20]	
Xiao ZY 2009	12	26	13	26	15.1%	0.86 [0.29, 2.55]	<del></del>
Yin LJ 2008	20	48	21	48	26.4%	0.92 [0.41, 2.06]	<del>-</del>
Zeng BZ 2010	2	31	4	30	8.2%	0.45 [0.08, 2.65]	-
Zeng CS 2014	10	30	12	30	17.3%	0.75 [0.26, 2.15]	<del></del>
Zheng C 2018	4	30	6	28	11.6%	0.56 [0.14, 2.26]	<del></del>
Total (95% CI)		195		195	100.0%	0.69 [0.44, 1.07]	•
Total events	54		69				
Heterogeneity: Chi2 =	1.98, df = 5	(P = 0.5)	85); I <sup>2</sup> = 0	%			
Test for overall effect:	Z = 1.65 (P	P = 0.10)	)				0.01 0.1 1 10 100 Favours [Control] Favours [Experimental]

C

	Experim	ental	Contr	ol		Odds Ratio	Odd	ls Ratio	
Study or Subgroup	Events	Total	<b>Events</b>	Total	Weight	M-H, Fixed, 95% C	I M-H, Fiz	xed, 95% CI	
Sun JH 2006	2	64	2	64	8.2%	1.00 [0.14, 7.33]		<del> </del>	
Wang HB 2013	19	43	21	42	50.0%	0.79 [0.34, 1.86]	_	_	
Xiong TQ 2010	1	26	1	26	4.1%	1.00 [0.06, 16.89]	-	+	
Zeng CS 2012	10	30	12	30	33.7%	0.75 [0.26, 2.15]		•	
Zhang SZ 2012	1	22	1	22	4.0%	1.00 [0.06, 17.07]	-	<del> </del>	
Total (95% CI)		185		184	100.0%	0.81 [0.45, 1.48]	<		
Total events	33		37						
Heterogeneity: Chi2 = 0	0.11, df = 4	(P = 1.0)	$(00); I^2 = 0$	%			0.01 0.1	1 10	100
Test for overall effect: 2	Z = 0.68 (P	= 0.49)						Favours [Exper	

D

	Experim	ental	Contr	ol		Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	<b>Events</b>	Total	Weight	M-H, Fixed, 95% C	M-H, Fixed, 95% CI
Sun JH 2006	5	64	5	64	50.2%	1.00 [0.27, 3.64]	<del></del>
Xiong TQ 2010	2	26	2	26	20.1%	1.00 [0.13, 7.69]	<del></del>
Zhang SZ 2012	2	22	3	22	29.7%	0.63 [0.10, 4.22]	
Total (95% CI)		112		112	100.0%	0.89 [0.35, 2.29]	•
Total events	9		10				
Heterogeneity: Chi2 = 0	0.17, df = 2	(P = 0.9)	92); I <sup>2</sup> = 0	%			0.01 0.1 1 10 100
Test for overall effect:	Z = 0.24 (P	= 0.81)					0.01 0.1 1 10 100 Favours [Control] Favours [Experimental]



_	Experim	ental	Contr	ol		Odds Ratio		Odds	Ratio		
Study or Subgroup	<b>Events</b>	Total	<b>Events</b>	Total	Weight	M-H, Fixed, 95% C		M-H, Fixe	ed, 95% C	:	
Wang HB 2013	6	43	7	42	68.5%	0.81 [0.25, 2.65]					
Zeng CS 2014	2	30	3	30	31.5%	0.64 [0.10, 4.15]					
Total (95% CI)		73		72	100.0%	0.76 [0.28, 2.06]		•	<b>-</b>		
Total events	8		10								
Heterogeneity: Chi <sup>2</sup> = (	0.04, df = 1	(P = 0.8)	84); I <sup>2</sup> = 0	%			0.01	0.1	-	10	100
Test for overall effect:	Z = 0.54 (P	= 0.59)						vours [Control]	Favours		

## $\mathbf{F}$

	Experime	ental	Contr	ol		<b>Odds Ratio</b>		Odds	Ratio	
Study or Subgroup	Events	Total	<b>Events</b>	Total	Weight	M-H, Fixed, 95% C		M-H, Fixe	ed, 95% CI	
Jiang CY 2013	11	30	24	33	75.7%	0.22 [0.07, 0.63]				
Zheng C 2018	3	30	5	28	24.3%	0.51 [0.11, 2.37]				
Total (95% CI)		60		61	100.0%	0.29 [0.12, 0.69]		•		
Total events	14		29							
Heterogeneity: Chi2 = 0	0.81, df = 1	(P = 0.3)	$37); I^2 = 0$	%			0.01	0.1	<del>                                     </del>	100
Test for overall effect:	Z = 2.79 (P	= 0.005	5)					ours [Control]		

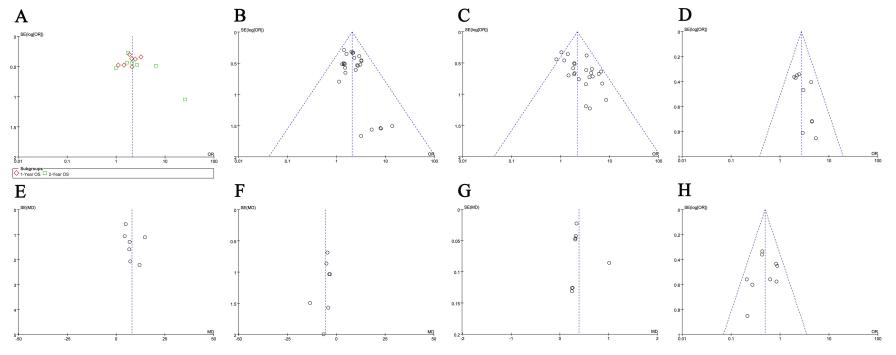
## G

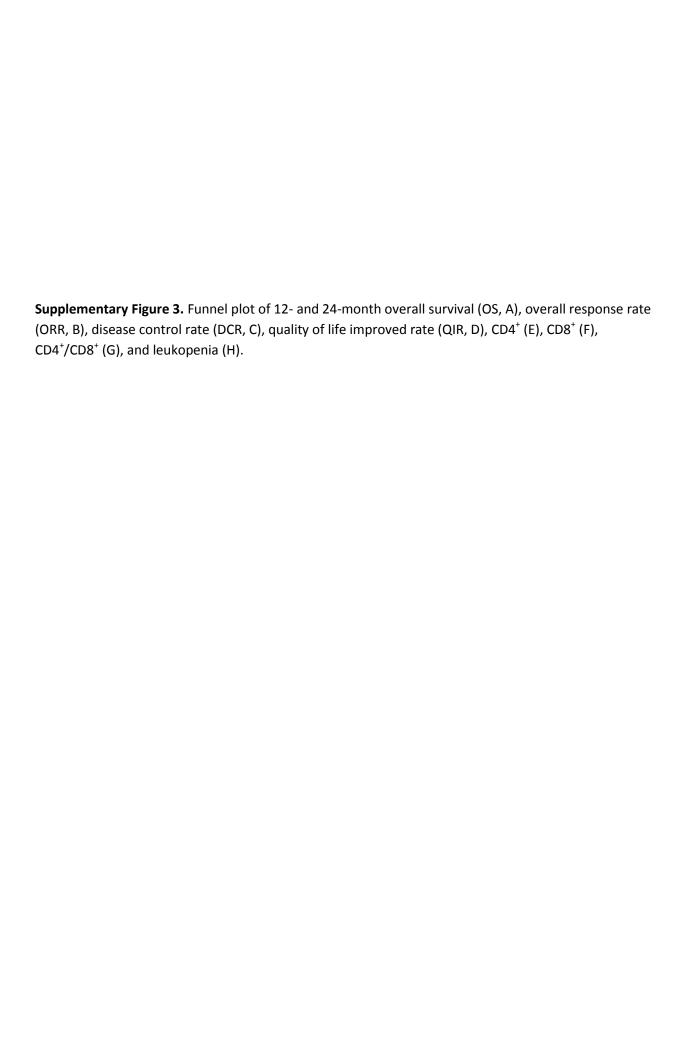
		Experime	ental	Contr	ol		Odds Ratio		Odds	Ratio		
Study	or Subgroup	Events	Total	<b>Events</b>	Total	Weight	M-H, Fixed, 95% C		M-H, Fixe	ed, 95% CI		_
Xie Yi	2003	6	31	15	31	49.0%	0.26 [0.08, 0.80]		_			
Zeng	BZ 2010	7	31	16	30	51.0%	0.26 [0.08, 0.77]		_			
Total	(95% CI)		62		61	100.0%	0.26 [0.12, 0.56]		•			
Total	events	13		31								
Heter	ogeneity: Chi <sup>2</sup> = 0	0.00, df = 1	(P = 1.0)	$00); I^2 = 0$	%			0.01	0.1	<del>                                     </del>	100	
Test f	or overall effect: 2	Z = 3.37 (P	= 0.000	07)					avours [Control]			al]

## H

	Experime	ental	Contr	ol		Odds Ratio		Odds	Ratio		
Study or Subgroup	Events	Total	<b>Events</b>	Total	Weight	M-H, Fixed, 95% C		M-H, Fixe	ed, 95% C		
Wang HB 2013	4	43	5	42	35.3%	0.76 [0.19, 3.05]			<del>                                     </del>		
Zeng CS 2014	9	30	12	30	64.7%	0.64 [0.22, 1.87]		_	$\vdash$		
Total (95% CI)		73		72	100.0%	0.68 [0.29, 1.60]			<b>-</b>		
Total events	13		17								
Heterogeneity: Chi2 = (	0.03, df = 1	(P = 0.8)	85); I <sup>2</sup> = 0	%			0.01	1	1 .	10	100
Test for overall effect: 2	Z = 0.88 (P	= 0.38)						0.1 rs [Control]	-		

**Supplementary Figure 2.** Forest plot of the comparison of adverse effects including leukopenia (A), gastrointestinal adverse effects (B), nausea and vomiting (C), anorexia (D), thrombocytopenia (E), hepatotoxicity (F), myelosuppression (G) and anemia (H) between the experimental and control group. Control group, conventional treatment alone group; Experimental group, conventional treatment and JLC combined group; JLC, Jinlong capsule. The fixed-effects meta-analysis model (Mantel-Haenszel method) was used.





Supplementary Table 1. Quality assessment of non-randomized comparative studies

Study			Non	-rando	mized	studie	es		Additio	Total			
	A	В	С	D	Е	F	G	Н	I	J	K	L	
Jia CH 2008 <sup>[19]</sup>	2	1	2	1	2	1	2	0	2	2	2	2	19
Li H 2007 <sup>[23]</sup>	2	1	2	1	1	2	2	0	2	2	2	2	19
Liang TJ 2005 <sup>[24]</sup>	2	1	2	1	2	2	2	0	2	2	2	2	20
Liu ZY 2015 <sup>[25]</sup>	2	1	2	1	2	1	2	0	2	2	2	2	19
Sun JH 2006 <sup>[27]</sup>	2	1	2	2	2	2	2	0	2	2	2	2	21
Wang HB 2013 <sup>[29]</sup>	2	1	2	2	2	2	2	0	2	2	2	2	21
Xiong TQ 2010 <sup>[34]</sup>	2	1	2	2	2	1	2	0	2	2	2	2	20
Zhang XQ 2012 <sup>[43]</sup>	2	1	2	2	2	2	2	0	2	2	2	2	21

**Notes:** The items are scored 0 (not reported), 1 (reported but inadequate) and 2 (reported and adequate).

A: A clearly stated aim; B: Inclusion of consecutive patients; C: Prospective collection of data; D: Endpoints appropriate to the aim of the study; E: Unbiased assessment of the study endpoint; F: Follow-up period appropriate to the aim of the study; G: Loss to follow up less than 5%; H: Prospective calculation of the study size; I: An adequate control group; J: Contemporary groups; K: Baseline equivalence of groups; L: Adequate statistical analyses.

**Supplementary Table 2.** Publication bias on therapeutic efficacy indexes (OS, ORR, DCR and QIR), immune function indexes (CD4<sup>+</sup>, CD8<sup>+</sup>, and CD4<sup>+</sup>/CD8<sup>+</sup>) and leucopenia.

<b>Publication Bias</b>		Therape	eutic efficacy in		Immı	ıne func	Adverse events		
	12-month OS	24-month OS	ORR	DCR	QIR	$CD4^{+}$	CD8 <sup>+</sup>	CD4 <sup>+</sup> /CD8 <sup>+</sup>	Leucopenia
Begg	0.764	0.133	0.168	0.035	0.107	0.133	0.548	0.902	0.675
Egger	0.301	0.161	0.049	0.001	0.036	0.321	0.331	0.603	0.542
Trim and fill analysis									
before			P < .0001	P < .0001	P < .0001				
after			P < .0001	P < .0001	P < .0001				

**Notes:** Parameters discussed in over 7 papers were conducted bias analyses.

Abbreviations: OS: overall survival; ORR: overall response rate; DCR: disease control rate; QIR: quality of life improved rate.

**Supplementary Table 3.** Subgroup analyses of ORR and DCR between the experimental and control group.

Parameter	Factors	Patients	Analysis	Hetero	geneity	Odds	95% CI	P-value
	at study level	Con/Exp	method	I <sup>2</sup> (%)	<i>P</i> -value	Ratio		
ORR	Therapeutic re	gimen						
	JLC+TACE	739/719	Fixed	0	0.96	1.93	1.56-2.38	< 0.00001
	JLC+SST	157/157	Fixed	0	0.97	7.37	1.91-28.36	0.004
	JLC+RT	117/116	Fixed	0	0.93	3.02	1.72-5.31	0.0001
	Study sample s	size						
	>80	719/683	Fixed	0	0.83	2.06	1.64-2.59	< 0.00001
	<80	379/380	Fixed	0	0.89	2.06	1.48-2.88	< 0.0001
	Type of contro	l trials						
	RCTs	677/654	Fixed	0	0.93	2.06	1.63-2.61	< 0.00001
	non-RCTs	421/409	Fixed	0	0.69	2.07	1.52-2.82	< 0.00001
DCR	Therapeutic re	gimen						
	JLC+TACE	672/653	Fixed	6	0.39	1.66	1.26-2.18	0.0003
	JLC+SST	157/157	Fixed	0	0.65	3.90	2.42-6.29	< 0.00001
	JLC+RT	117/116	Fixed	0	0.76	4.50	1.46-13.86	0.009
	Study sample s	size						
	>80	652/617	Fixed	24	0.23	1.62	1.22-2.15	0.0009
	<80	379/380	Fixed	0	0.91	3.46	2.40-4.99	< 0.00001
	Type of contro	l trials						
	RCTs	610/588	Fixed	0	0.83	2.48	1.84-3.33	< 0.00001
	non-RCTs	421/409	Random	63	0.01	2.09	1.11-3.94	0.02

**Notes:** Con, Control group (Conventional treatment alone group); Exp, Experimental group (Conventional treatment and JLC combined group).

**Abbreviations:** ORR: overall response rate; DCR: disease control rate; JLC: Jinlong capsule; TACE: transcatheter arterial chemoembolization; SST: Support and symptomatic treatment; RT: Radiotherapy; RCTs; randomized controlled trials.