

Efficacy of the No. 10 lymphadenectomy with spleen preservation on patients with gastric cancer and/or esophagogastric junction adenocarcinoma who underwent total gastrectomy: a systematic review and meta-analysis

Bo-Wei Xia¹, Chen Wang¹, Yong-Yong Liu¹, Yong Fan¹, Xiao-Dong He¹, Ying-Xin Kang¹, Xin-Yuan Zhou¹, Xiao-Lu Su², Yue-Bin Wang¹, Min-Xue Chen¹, Bo-Xiong Kang¹

¹Department of General Surgery, Lanzhou University Second Hospital, Lanzhou, China; ²Department of Pathology, Lanzhou University Second Hospital, Lanzhou, China

Contributions: (I) Conception and design: BW Xia, BX Kang; (II) Administrative support: C Wang, XD He; (III) Provision of study materials or patients: MX Chen, BW Xia; (IV) Collection and assembly of data: YB Wang, XL Su, XY Zhou; (V) Data analysis and interpretation: MX Chen, Y Fan, YY Liu, YX Kang; (VI) Manuscript writing: All authors; (VII) Final approval of manuscript: All authors.

Correspondence to: Bo-Xiong Kang. Doctor, Professor, Department of General Surgery, Lanzhou University Second Hospital, No. 82, Cuiying Road, Lanzhou 730030, China. Email: kangbx2000@163.com.

Background: Surgery with total gastrectomy and D2 lymph node dissection (LND) has been recommended as the standard treatment for patients with advanced upper and middle gastric carcinoma and/ or Siewert type II/III adenocarcinoma of the esophagogastric junction (AEG). However, whether the No. 10 lymph node (No. 10 LN, also known as splenic hilar LN) should be dissected in total gastrectomy remains controversial. We aimed to evaluate whether the No. 10 LND with spleen preservation has survival benefit for patients with gastric cancer and/or AEG who underwent the total gastrectomy.

Methods: The PubMed, Embase, the Cochrane Library, ClinicalTrials.gov and American Society of Clinical Oncology.org (ASCO.org) were electronically searched to identify eligible studies. The primary outcome was the survival rate, and secondary outcomes included the disease-free survival (DFS) rate and side effects. The Review Manager 5.3.5 software was used for the meta-analysis. The odds ratio (OR) and mean difference with 95% confidence interval (CI) were calculated. The statistical heterogeneity was assessed using chi-square (χ^2) and I² tests.

Results: Eight studies enrolling a total of 4,131 patients were eligible for our review. The meta-analysis results demonstrated that the No. 10 LND group was significantly better than the non-No. 10 LND group in terms of the 3- (OR =0.71, 95% CI: 0.62–0.81, P<0.00001) and the 5-year (OR =0.66, 95% CI: 0.58–0.75, P<0.00001) survival rates but not in the 1-year survival rate (OR =0.91, 95% CI: 0.75–1.11, P=0.36). The DFS rates in the No. 10 LND group were significantly increased after 1 (OR =0.76, 95% CI: 0.61–0.93, P=0.008), 3 (OR =0.69, 95% CI: 0.60–0.81, P<0.00001), and 5 (OR =0.66, 95% CI: 0.56–0.76, P<0.00001) years compared with those in the non-No. 10 LND group.

Discussion: Evidence shows that the No. 10 LND with spleen preservation can improve the survival and the DFS rates for patients with gastric cancer and/or Siewert type II/III AEG who underwent the total gastrectomy. High-quality prospective trials are expected.

Keywords: No. 10 lymphadenectomy; gastric cancer; adenocarcinoma of the esophagogastric junction (AEG); total gastrectomy; systematic review

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Introduction

Gastric cancer is responsible for over one million new cases in 2020, and is listed as the fourth most common malignancy and one of the most common causes of cancerrelated death worldwide (1,2). According to the Japanese Gastric Cancer Treatment Guideline (JGCTG) 2010 (version 3), surgery with total gastrectomy and D2 lymph node dissection (LND) is recommended as the standard treatment for patients with advanced upper and middle gastric carcinoma in East Asia (3). The advanced Siewert type II/III adenocarcinoma of the esophagogastric junction (AEG) is suggested with the same treatment with proximal gastric cancer as their anatomical position and biological behavior are quite similar (3).

Nevertheless, whether the No. 10 lymph node (No.10 LN, also known as splenic hilar LN) should be dissected in total gastrectomy remains controversial. The incidence rate of No. 10 lymph node metastasis (LNM) is reported to be 9.0–27.9% in the advanced upper and middle gastric cancers (4-7) and 4.8–15.0% in the Siewert type II/III AEG (8-11). Thus, the No. 10 LND is recommended to be added in the total gastrectomy (12).

However, some studies consider that undergoing the No. 10 LND is unnecessary, for the No. 10 LNM is considered as one of the incurable factors of prognosis and the No. 10 LND does not have a survival benefit (13,14). Shin *et al.* (15) have reported that patients with the No. 10 LNM have worse survival than patients without the No. 10 LNM and does not gain survival benefit from the No. 10 LND. Recently, the No. 10 LN has been deleted from the definition of D2 LND in total gastrectomy according to the JGCTG 2018 (5th edition) by the Japanese Gastric Cancer Association (16).

The aim of the present study was to perform a systematic review of randomized controlled trials (RCTs) or cohort studies to evaluate the effects of No. 10 LND with spleen preservation on patients with gastric cancer and/or Siewert type II/III AEG who have undergone total gastrectomy. We present the following article in accordance with the PRISMA reporting checklist (17) (available at https://tcr. amegroups.com/article/view/10.21037/tcr-22-522/rc).

Methods

Search strategy and selection criteria

The PubMed, Embase, the Cochrane Library, ClinicalTrials.gov and American Society of Clinical Oncology.org (ASCO.org) were searched until January 2022 for relevant citations. A combination of the following terms was used to complete the search: "gastric cancer", "cardiac carcinoma", "esophagogastric junction carcinoma", "No. 10 lymphadenectomy", "splenic hilar lymphadenectomy", "No. 10 lymph node", "splenic hilar lymph node", and "gastrectomy". Manual searches included scanning of reference lists in relevant articles. No language restriction was applied.

Eligible trials were RCTs or cohort studies that compared the No. 10 LND with the non-No.10 LND for patients with gastric cancer and/or Siewert type II/III AEG who underwent the total gastrectomy. Moreover, data concerning the survival rate should be reported in studies. Studies were excluded if they met the following criteria: (I) intraoperative evidence of peritoneal dissemination or distant metastasis; (II) combined major organ resection (except necessary organ resection, such as splenectomy or pancreatectomy, because of intraoperative organ injury, the intraoperative detection of invasion of the pancreas or spleen, or to enable the en bloc dissection of evident metastatic No. 10 LNs); (III) incomplete pathological data; (IV) and neoadjuvant therapy.

Study selection

Two authors (MXC, BWX) independently selected literature. A third author (BXK) resolved any discrepancies if the first two authors disagreed. The full text of each potentially eligible study was evaluated for inclusion or exclusion in accordance with the selection criteria of the two independent reviewers (YF and YXK).

Data extraction

Two authors (XLS and YBW) independently extracted the data using predefined data extraction forms. Extracted data included study details, study population characteristics, interventions, and outcomes from each eligible trial. All relevant texts, tables, and figures were reviewed for data extraction. Any disagreement in data extraction was resolved by a third reviewer (YYL).

Quality assessment

The methodological quality of cohort studies was assessed by using of the Newcastle-Ottawa Scale (18). A "star system" was applied in each study in accordance with three broad perspectives: selection of cases (0–4 stars), comparability



Figure 1 Flow diagram summarizing study identification and selection. ASCO, American Society of Clinical Oncology; RCT, randomized controlled trials.

of groups (0-2 stars), and assessment of outcome (0-3 stars). The quality of each study was graded as either "high quality" (8-9 stars) or "moderate quality" (5-7 stars). The quality of the included RCTs was assessed using the modified Jadad standard. Three items were included in the specified criteria of the RCTs: randomization (0-2 points), dropout or withdrawal (0-1 point), and allocation concealment (0-2 points) (19). A blind approach was discarded in RCTs because of the intrinsic nature of the intervention. Studies that received an Jadad score of 3 or higher were classified as highquality studies. Two authors (YYL and YXK) independently ranked and assessed each study.

Outcome measures

The primary outcome was overall survival rate that was assessed in the included study. This endpoint was measured over one, three, or five years, depending on the size of the study and the length of the follow-up. The diseasefree survival (DFS) rate at each time point, complications and mortality associated with the No. 10 LND were the secondary outcomes.

Statistical analysis

A meta-analysis of outcomes by combining various studies

was performed using the Review Manager (RevMan) software, version 5.3.5 (The Cochrane Collaboration, Software Update, Oxford). The effect measures of interest were odds ratio (OR) and mean difference with 95% confidence interval (CI). The statistical heterogeneity was assessed using chi-square (χ^2) and I² tests. I² <25%, 25% \leq $I^2 \leq 50\%$, and $I^2 > 50\%$ reflected small, moderate, and large inconsistencies, respectively. Sub-group analyses or sensitivity analyses was undertaken to attempt to explain heterogeneities if existed. The ability to conduct subgroup analyses depended on whether the required information was reported in the included studies. The location of tumor was considered for possible subgroup analysis. Sensitivity analyses were carried out only in high quality trials to avoid errors caused by poor quality studies. The publication bias was performed using the Begg's funnel plots and Egger's tests if the number of included studies was more than 10 (20,21). P<0.05 was considered significant.

Results

Selection of trials

From the 656 citations identified using database searches, 18 duplicate studies and 623 reviews, case reports, letters, editorials, and irrelevant articles were excluded (Figure 1).

Furthermore, 7 out of the 15 remaining trials with potential relevance were excluded from the study after reading the full text. One study referred to the posterior No. 10 LND (22); and six studies did not reveal sufficient outcome information to perform analysis (23-28). Finally, eight retrospective cohort studies involving 4,131 patients were included in the study (14,29-35).

Among the included studies, 1,929 patients were in the No. 10+ group (No. 10 LND was conducted for patients with gastric cancer and/or Siewert type II/III AEG who underwent the total gastrectomy) and 2,202 patients were in the No. 10- group (No. 10 LND was not conducted for patients with gastric cancer and/or Siewert type II/III AEG who underwent total gastrectomy). For the study by Park et al. (14), only the data of arms fulfilled the included criteria and were extracted for meta-analysis. In the study of Bian et al. (30), although the data fulfilled the included criteria was limited in the patients with negative No. 4s LNs, we performed a sensitivity analysis and found the result be influenced slightly by the confinement of patients with negative No. 4s LNs. Thus, we included this study for meta-analysis. The average age of patients was 58.9 years, and 74.8% of the patients were male (Table 1). The percentages of TNM stages I-IV of the tumors were 11.0%, 25.7%, 59.5%, and 3.8%, respectively, for patients with the No. 10 LND, and 16.1%, 25.9%, 54.3%, and 3.7%, respectively, for patients without the No. 10 LND. The mean tumor size of gastric cancer and/or Siewert type II/III AEG was 5.9 cm. The mean duration of follow-up was 70.7 months.

Qualitative analysis of studies

As *Table 2* shows, among the eight cohort studies, 7 were of high quality (14,29,30,32-35), and 1 was of moderate quality (31).

Survival rate

Overall survival rate

The meta-analysis results showed that the No. 10 LND significantly improved the 3- [eight studies (14,29-35) reported this data, OR =0.71, 95% CI: 0.62–0.81, P<0.00001; heterogeneity: I^2 =39%, P=0.12 for χ^2] and 5-year [eight studies (14,29-35) reported this data, OR =0.66, 95% CI: 0.58–0.75, P<0.00001; heterogeneity: I^2 =24%, P=0.23 for χ^2] survival rates but not the 1-year survival rate [eight studies (14,29-35) reported this data, OR =0.91, 95% CI: 0.75–1.11, P=0.36; heterogeneity: I^2 =53%, P=0.04 for χ^2 ; *Figure 2*].

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Survival rate of patients with gastric cancer after surgical resection with total gastrectomy

The pooled meta-analysis results demonstrated that the No. 10 LND significantly improved the 3- [six studies (14,30,32-35) reported this data, OR =0.76, 95% CI: 0.65–0.90, P=0.001; heterogeneity: $I^2 = 32\%$, P=0.19 for χ^2] and 5-year [seven studies (14,29,30,32-35) reported this data, OR =0.70, 95% CI: 0.60–0.82, P<0.00001; heterogeneity: $I^2 = 13\%$, P=0.33 for χ^2] survival rates but not the 1-year survival rate [six studies (14,30,32-35) reported this data, OR =0.95, 95% CI: 0.74–1.21, P=0.67; heterogeneity: $I^2 = 61\%$, P=0.02 for χ^2 ; *Figure 3A*].

Survival rate of patients with gastric cancer and type III AEG after surgical resection with total gastrectomy

The overall survival rates in patients with the No. 10 LND were significantly higher than those in patients without the No. 10 LND after 3 [seven studies (14,30-35) reported this data, OR =0.72, 95% CI: 0.57–0.90, P=0.004; heterogeneity: I² =54%, P=0.04 for χ^2], and 5 years [eight studies (14,29-35) reported this data, OR =0.64, 95% CI: 0.53–0.79, P<0.0001; heterogeneity: I² =49%, P=0.06 for χ^2], but not 1 year [seven studies (14,30-35) reported this data, OR =0.92, 95% CI: 0.62–1.37, P=0.69; heterogeneity: I² =63%, P=0.01 for χ^2 ; *Figure 3B*].

DFS rate

Overall DFS rate

The meta-analysis showed significant differences between the two groups, and the result favored the No. 10+ group with 1- [five studies (30-33,35) reported this data, OR =0.76, 95% CI: 0.61–0.93, P=0.008; heterogeneity: I^2 =0, P=0.61 for χ^2], 3- [five studies (30-33,35) reported this data, OR =0.69, 95% CI: 0.60–0.81, P<0.00001; heterogeneity: I^2 =0%, P=0.55 for χ^2], and 5-year [five studies (30-33,35) reported this data, OR =0.66, 95% CI: 0.56–0.76, P<0.00001; heterogeneity: I^2 =5%, P=0.38 for χ^2] DFS rates (*Figure 4*).

DFS rate of patients with gastric cancer after surgical resection with total gastrectomy

Results showed that the No. 10 LND was associated with a significant improvement in the 1- [four studies (30,32,33,35) reported this data, OR =0.79, 95% CI: 0.62–0.99, P=0.04; heterogeneity: $I^2 = 0\%$, P=0.54 for χ^2], 3- [four studies (30,32,33,35) reported this data, OR =0.72, 95% CI: 0.60–0.85, P=0.0002; heterogeneity: $I^2 = 0\%$, P=0.47 for χ^2], and

Table 1	Basic cl	naracterist	tics of the include	ed studies								
Study	Year	Country	Study type	Arms	Sample size (n)	Sex, M/F (n)	Age (year)	Tumor size (cm)	Tumor location (upper third and AEG/middle third/lower third/entire and linitisplastica)	Pathological differentiation (differentiated/ undifferentiated)	TNM stage [†] (I/II/II/IV)	Follow-up duration (month)
Bian	2016	China	Retrospective	No. 10+ [‡]	260	185/75	NC	NC	NC	39/221	5/59/196/0	62
				No. 10 ^{-§}	243	170/73	NC	NC	NC	48/195	6/58/179/0	62
Huang	2017	China	Retrospective	No. 10+	198	155/43	61.9	6.1	80/75/0/43	NC	9/49/140/0	60
				No. 10–	198	154/44	61.1	5.9	83/80/0/35	NC	15/45/138/0	60
Lin	2021	China	Retrospective	No. 10+	354	280/74	62.3	5.3	161/122/0/71	275/79	61/90/203/0	60
				No. 10–	354	286/68	63.1	5.3	155/146/0/53	289/65	73/85/196/0	60
Liu	2021	China	Retrospective	No. 10+	237	165/72	58.0	6.8	148/60/19/10	68/169	20/44/156/17	91.2
				No. 10–	237	172/65	58.1	7.0	138/64/22/13	68/169	20/34/163/20	91.2
۲۷	2016	China	Retrospective	No. 10+	293	241/52	NC	4.9	293/0/0/0	NC	52/87/154/0	47
				No. 10–	401	335/66	NC	4.9	401/0/0/0	NC	81/104/216/0	47
hO	2021	Korea	Retrospective	No. 10+	288	194/94	56.1	7.5	NC	88/200	NC	60
				No. 10–	288	198/90	56.1	7.5	NC	89/199	NC	60
Park	2019	Korea	Retrospective	No. 10+	79	58/21	55.5	NC	NC	NC	22/57/0/0	93.3
				No. 10–	248	168/80	55.1	NC	NC	NC	100/148/0/0	93.3
Yang	2014	China	Retrospective	No. 10+	220	163/57	NC	NC	104/64/41/11	35/185	12/35/128/45	89.5
				No. 10–	233	166/67	NC	NC	64/85/51/33	25/208	14/22/146/51	96
[†] , TNM s dissectic	stage w n. M/F,	as accor male/fer	ding to seventh nale; AEG, ader	American Jc tocarcinoma	bint Commi a of the esc	ittee on Car pphagogastr	ncer stag ric juncti	jing system on; TNM, tu	for gastric cancer; [‡] , No. 1 mor-node-metastasis; NC,	0 lymph node dissect not clear.	tion; [§] , non-No. 10	ymph node

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Table 2 Quality assessment of included cohort studies

Chudu		Sele	ction		Compa	arability		Outcome		Coore
Sludy	1	2	3	4	5A	5B	6	7	8	Score
Bian 2016	*†	*	*	*	*	*	*	*	*	******
Huang 2017	*	*	*	*	*	*	*	*	*	******
Lin 2021	*	*	*	*	*	*	*	*	*	******
Liu 2021	*	*	*	*	*	*	*	*	*	******
Lv 2016	*	*	*	*			*	*	*	*****
Oh 2021	*	*	*	*	*	*	*	*	*	*****
Park 2019	*	*	*	*	*		*	*	*	*****
Yang 2014	*	*	*	*	*	*	*	*	*	******

1, representativeness of the exposed cohort; 2, non-exposed cohort drawn from the same community; 3, ascertainment of exposure; 4, outcome of interest not present at start of study; 5A, comparability of cohorts on the basis of tumor categories; 5B, comparability of cohorts on the other factors; 6, assessment of outcomes; 7, follow-up long enough for outcomes to occur; 8, adequacy of follow up of cohorts; [†], each star represents if an individual criterion within the subsection was fulfilled; [‡], score for study quality. The quality of each included cohort study was graded as either "high quality" (8–9 stars) or "moderate quality" (5–7 stars).

	No. 1	0+	No.	10–		Odds ratio (Non-eve	ent) Odds ratio (Non-event)
Study or subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% C	M-H, Fixed, 95% Cl
1.1.1 1-year survival							
Bian 2016	231	260	231	243	5.3%	2.42 [1.20, 4.85]	
Huang 2017	174	198	171	198	11.4%	0.87 [0.48, 1.57]	
Lin 2021	321	354	303	354	22.2%	0.61 (0.38, 0.97)	_
Liu 2021	204	237	195	237	17.4%	0.75 [0.46, 1.23]	
Lv 2016	268	293	353	401	17.8%	0.69 [0.41, 1.14]	
Oh 2021	263	288	264	288	10.5%	1.05 (0.58, 1.88)	_
Park 2019	74	79	240	248	1.7%	2.03 [0.64, 6.38]	
Yang 2014	184	220	198	233	13.7%	1.11 [0.67, 1.84]	
Subtotal (95% CI)		1929		2202	100.0%	0.91 [0.75, 1.11]	•
Total events	1719		1955				
Heterogeneity: Chi ² =	14.81 df=	= 7 (P =	0.04): P	= 53%			
Test for overall effect	7 = 0.91 (P=03	6)				
			-,				
1.1.2 3-year survival							
Bian 2016	164	260	138	243	131%	0 77 10 54 1 101	
Huang 2017	129	198	109	198	11 1 %	0.66 (0.44 0.98)	
L in 2021	253	354	216	354	18.9%	0.62/0.46/0.861	_ _
Liu 2021	140	237	117	237	13.6%	0.68 [0.47, 0.97]	
Lv 2016	229	293	262	401	17.5%	0.53 (0.37, 0.74)	_
Oh 2021	204	288	198	288	12.2%	0.91 [0.63, 1.29]	
Park 2019	68	79	227	248	1.7%	1,75 (0.80, 3,81)	
Yang 2014	118	220	113	233	12.0%	0.81 [0.56, 1.18]	
Subtotal (95% CI)		1929		2202	100.0%	0.71 [0.62, 0.81]	•
Total events	1305		1380				
Heterogeneity: Chi ² =	11.40. df=	= 7 (P =	= 0.12); P	² = 39%			
Test for overall effect:	Z = 5.02 (P < 0.0	0001)				
1.1.3 5-year survival							
Bian 2016	134	260	104	243	12.8%	0.70 [0.49, 1.00]	
Huang 2017	123	198	94	198	11.2%	0.55 [0.37, 0.82]	
Lin 2021	225	354	185	354	18.6%	0.63 [0.46, 0.85]	
Liu 2021	121	237	84	237	13.5%	0.53 [0.36, 0.76]	
Lv 2016	210	293	241	401	16.7%	0.60 [0.43, 0.82]	
Oh 2021	179	288	165	288	13.2%	0.82 [0.59, 1.14]	
Park 2019	63	79	209	248	2.6%	1.36 [0.71, 2.60]	
Yang 2014	103	220	87	233	11.5%	0.68 [0.47, 0.98]	
Subtotal (95% CI)		1929		2202	100.0%	0.66 [0.58, 0.75]	◆
Total events	1158		1169				
Heterogeneity: Chi ² =	9.26, df =	7 (P =	0.23); I ² :	= 24%			
Test for overall effect:	Z = 6.37 (P < 0.0	0001)				
							Favours No. 10+ Favours No. 10-

Figure 2 Meta-analysis of overall survival in trials comparing No. 10 LND versus non-No. 10 LND for patients with gastric cancer and/or Siewert type II/III AEG who have undergone total gastrectomy. No. 10 LND, No. 10 lymph node dissection; AEG, adenocarcinoma of the esophagogastric junction; No. 10+, No. 10 lymph node dissection group; No. 10–, non-No. 10 lymph node dissection group; M-H, Mantel-Haenszel; CI, confidence interval.



Figure 3 Meta-analysis of survival in trials comparing No. 10 LND versus non-No. 10 LND for: (A) patients with gastric cancer who have undergone total gastrectomy; (B) patients with gastric cancer and/or Siewert type III AEG who have undergone total gastrectomy. No. 10 LND, No. 10 lymph node dissection; AEG, adenocarcinoma of the esophagogastric junction; No. 10+, No. 10 lymph node dissection group; No. 10-, non-No. 10 lymph node dissection group; M-H, Mantel-Haenszel; CI, confidence interval.

	No. 1	0+	No. 1	0-		Odds ratio (Non-ever	nt) Odds ratio (Non-event)
Study or subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% (CI M-H, Random, 95% CI
1.4.1 1-year disease-	free surv	ival					
Bian 2016	233	260	209	243	14.8%	0.71 [0.42, 1.22]	
Huang 2017	163	198	153	198	17.6%	0.73 [0.45, 1.20]	
Lin 2021	307	354	290	354	25.6%	0.69 [0.46, 1.04]	
Lv 2016	259	293	334	401	21.8%	0.65 [0.42, 1.02]	
Oh 2021	244	288	246	288	20.4%	1.06 [0.67, 1.67]	
Subtotal (95% CI)		1393		1484	100.0%	0.76 [0.61, 0.93]	•
Total events	1206		1232				
Heterogeneity: Tau ² =	0.00; Chi	² = 2.69	0, df = 4 (P = 0.6	1); $l^2 = 0\%$		
Test for overall effect:	Z = 2.65 (P = 0.0	08)				
1.4.2 3-year disease-	free surv	ival					
Bian 2016	149	260	129	243	18.8%	0.84 [0.59, 1.20]	
Huang 2017	129	198	108	198	14.2%	0.64 [0.43, 0.96]	
Lin 2021	244	354	204	354	24.5%	0.61 [0.45, 0.83]	
Lv 2016	211	293	247	401	22.1%	0.62 [0.45, 0.86]	
Oh 2021	187	288	172	288	20.4%	0.80 [0.57, 1.12]	
Subtotal (95% CI)		1393		1484	100.0%	0.69 [0.60, 0.81]	◆
Total events	920		860				
Heterogeneity: Tau ² =	0.00; Chi	² = 3.05	5, $df = 4$ (P = 0.5	5); I ² = 0%		
Test for overall effect:	Z=4.68 (P < 0.0	0001)				
1.4.3 5-year disease-	free surv	ival					
Bian 2016	123	260	96	243	18.1%	0.73 [0.51, 1.04]	
Huang 2017	123	198	94	198	14.3%	0.55 [0.37, 0.82]	
Lin 2021	214	354	184	354	25.2%	0.71 [0.53, 0.95]	
Lv 2016	211	293	230	401	21.6%	0.52 [0.38, 0.72]	
Oh 2021	171	288	153	288	20.8%	0.78 [0.56, 1.08]	
Subtotal (95% CI)		1393		1484	100.0%	0.66 [0.56, 0.76]	•
Total events	842		757				
Heterogeneity: Tau ² =	0.00; Chi	² = 4.20), df = 4 (P = 0.3	8); l² = 5%		
Test for overall effect:	Z= 5.38 (P < 0.0	0001)				
						-	0.2 0.5 1 2 5
							Favours No. 10+ Favours No. 10-

Figure 4 Meta-analysis of overall DFS in trials comparing No. 10 LND versus non-No. 10 LND for patients with gastric cancer and/or Siewert type II/III AEG who have undergone total gastrectomy. DFS, disease-free survival; No. 10 LND, No. 10 lymph node dissection; AEG, adenocarcinoma of the esophagogastric junction; No. 10+, No. 10 lymph node dissection group; No. 10-, non-No. 10 lymph node dissection group; M-H, Mantel-Haenszel; CI, confidence interval.

5-year [four studies (30,32,33,35) reported this data, OR =0.70, 95% CI: 0.59–0.83, P<0.0001; heterogeneity: $I^2 = 0\%$, P=0.62 for χ^2] DFS rates (*Figure 5A*).

DFS rate of patients with gastric cancer and type III AEG after surgical resection with total gastrectomy

The DFS rate in the No. 10+ group was significantly higher after 1 [five studies (30-33,35) reported this data, OR =0.72, 95% CI: 0.55–0.93, P=0.01; heterogeneity: I^2 =30%, P=0.22 for χ^2], 3 [five studies (30-33,35) reported this data, OR =0.66, 95% CI: 0.53–0.82, P=0.0001; heterogeneity: I^2 =43%, P=0.14 for χ^2], and 5 years [five studies (30-33,35) reported this data, OR =0.61, 95% CI: 0.46–0.80, P=0.0004; heterogeneity: I^2 =67%, P=0.02 for χ^2] than that in the No. 10– group (*Figure 5B*).

Safety

The most frequent side effects correlated with the No. 10 LND that were reported in the trials were iatrogenic spleen injury, intraoperative blood loss, pancreas-related complications, and peritoneal bleeding. The pooled metaanalysis results demonstrated that there was no significant difference between the two groups in the complications with grade I-II [five studies (31-35) reported this data, OR =1.20, 95% CI: 0.96–1.51, P=0.12; heterogeneity: I² =10%, P=0.35 for χ^2], complications with grade III-IV [five studies (31-35) reported this data, OR =1.30, 95% CI: 0.91–1.85, P=0.16; heterogeneity: I² =21%, P=0.28 for χ^2], and mortality [five studies (29,30,32-34) reported this data, OR =1.59, 95% CI: 0.52–4.87, P=0.42; heterogeneity: I² =0%, P=0.72 for χ^2 , *Figure 6*].

Discussion

This systematic review shows that the No. 10 LND with spleen preservation can significantly improve the overall survival and the DFS rates of patients with gastric cancer and/or Siewert type II/III AEG who have undergone the total gastrectomy.

Given the special anatomical position of the spleen and various complicated splenic hilum vessels, the exposure and the vascularization of splenic vessels are difficult to perform. Moreover, pancreas-related complications and bleeding

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Α		No.	10+	No.	10-		Odds ratio (Non-event)	Odds ratio (Non-event)
ŕ ,	Study or subgroup	Event	s Total	Event	s Tota	I Weight	M-H, Random, 95% Cl	M-H, Random, 95% Cl
	1.5.1 1-year disease-fr	ee surv	ival					
	Bian 2016	233	260	209	243	18.9%	0.71 [0.42, 1.22]	
	Huang 2017	163	198	153	198	22.4%	0.73 [0.45, 1.20]	
	Lin 2021	307	354	290	354	32.7%	0.69 [0.46, 1.04]	
	Subtotal (95% CI)	244	1100	240	1083	100.0%	0.79 [0.62, 0.99]	•
	Total events	947		898	1000	1001070	0110 [0102, 0100]	
	Heterogeneity: Tau ² = 0	0.00: Chi	² = 2.17	df = 3 (P = 0.5	4): $ ^2 = 0\%$		
	Test for overall effect: Z	= 2.01 ((P = 0.04	4)				
	1.5.2 3-year disease-fr	ee surv	ival			0.000		
	Bian 2016	149	260	129	243	24.1%	0.84 [0.59, 1.20]	
	Huang 2017	129	198	108	198	18.3%	0.64 [0.43, 0.96]	
	Lin 2021	244	354	204	354	31.4%	0.61 [0.45, 0.83]	
	Subtotal (95% CI)	187	288	172	288	20.2%	0.72 [0.60, 0.85]	•
	Total events	709	1100	613	1005	100.070	0.12 [0.00, 0.03]	•
	Heterogeneity: Tau ² = 0).00: Chi	² = 2.50	. df = 3 (P = 0.4	7): I ² = 0%		
	Test for overall effect: Z	= 3.79 ((P = 0.00	002)				
	1.5.3 5-year disease-fr	ee surv	ival					
	Bian 2016	123	260	96	243	23.0%	0.73 [0.51, 1.04]	
	Huang 2017	123	198	94	198	18.0%	0.55 [0.37, 0.82]	
	Lin 2021	214	354	184	354	32.5%	0.71 [0.53, 0.95]	
	Subtotal (95% CI)	171	1100	155	1083	100.0%	0.78 [0.56, 1.06]	•
	Total events	631		527		1001010		
	Heterogeneity: Tau ² = 0	0.00; Chi	P = 1.79	, df = 3 (P = 0.6	2); I² = 0%		
	Testion overall ellect. Z	- 4.151	(F < 0.00	001)				
							0.2	0.5 1 2 5
							Fav	ours No. 10+ Favours No. 10-
_		N		N				
В	o , 1	_NO.	10+	NO.	10-		Odds ratio (Non-event)	Odds ratio (Non-event)
	Study or subgroup	Event	s lotal	Event	s lota	i weight	M-H, Random, 95% CI	M-H, Random, 95% Cl
	Rian 2016	233	260	200	243	17 3%	0 71 10 42 1 221	
	Huang 2017	163	198	153	198	19.6%	0.73 [0.45 1.20]	
	Lin 2021	307	354	290	354	25.2%	0.69 [0.46, 1.04]	
	Lv 2016	155	175	166	215	16.1%	0.44 [0.25, 0.77]	
	Oh 2021	244	288	246	288	21.7%	1.06 [0.67, 1.67]	
	Subtotal (95% CI)		1275		1298	100.0%	0.72 [0.55, 0.93]	•
	Total events	1102		1064				
	Test for overall effect: Z	.03; Chi = 2.51 (P = 0.0	l,αt=4 (1)	P = 0.2	2); 1* = 30%		
	1.6.2 3-year disease-fr	ee surv	ival					
	Bian 2016	149	260	129	243	20.6%	0.84 [0.59, 1.20]	
	Huang 2017	129	198	108	198	17.5%	0.64 [0.43, 0.96]	
	Lin 2021	244	354	204	354	23.7%	0.61 [0.45, 0.83]	
	Lv 2016	122	175	108	215	16.7%	0.44 [0.29, 0.67]	
	Oh 2021	187	288	172	288	21.6%	0.80 [0.57, 1.12]	
	Subtotal (95% CI)	001	1275	704	1298	100.0%	0.66 [0.53, 0.82]	-
	Total events	8.51		121				
	Heterogeneity: Tau ² = 0 Test for overall effect: Z).03; Chi = 3.80 (P = 0.00	l, df = 4 (001)	(P = 0.1	4); I² = 43%		
	Heterogeneity: Tau ² = 0 Test for overall effect: Z).03; Chi := 3.80 (P = 7.00 P = 0.01	l, df = 4 (001)	P = 0.1	4); I ² = 43%		
	Heterogeneity: Tau ² = 0 Test for overall effect: Z 1.6.3 5-year disease-fr Bian 2016).03; Chi := 3.80 (ree surv 123	P = 7.00 P = 0.01 ival 260	1, df = 4 (001) 96	P = 0.1	4); I² = 43% 20.2%	0.73 (0.51 1 04)	_
	Heterogeneity: Tau ² = 0 Test for overall effect: Z 1.6.3 5-year disease-fr Bian 2016 Huang 2017	0.03; Chi = 3.80 (ree surv 123 123	P = 7.00 P = 0.00 ival 260 198	1, df = 4 (001) 96 94	P = 0.1 243 198	4); I ² = 43% 20.2% 18.5%	0.73 [0.51, 1.04] 0.55 [0.37. 0.82]	
	Heterogeneity: Tau ² = 0 Test for overall effect: Z 1.6.3 5-year disease-fr Bian 2016 Huang 2017 Lin 2021	0.03; Chi = 3.80 (ree surv 123 123 214	ival 260 198 354	I, df = 4 (001) 96 94 184	P = 0.1 243 198 354	4); I ² = 43% 20.2% 18.5% 22.3%	0.73 [0.51, 1.04] 0.55 [0.37, 0.82] 0.71 [0.53, 0.95]	
	Heterogeneith; Tau ² = 0 Test for overall effect: Z 1.6.3 5-year disease-fr Bian 2016 Huang 2017 Lin 2021 Lv 2016	0.03; Chi = 3.80 (ree surv 123 123 214 122	ival 260 198 354 175	1, df = 4 (001) 96 94 184 93	243 198 354 215	4); I ² = 43% 20.2% 18.5% 22.3% 17.8%	0.73 [0.51, 1.04] 0.55 [0.37, 0.82] 0.71 [0.53, 0.95] 0.33 [0.22, 0.50]	
	Heterogeneity: Tau" = 0 Test for overall effect: Z 1.6.3 5-year disease-fr Bian 2016 Huang 2017 Lin 2021 Lv 2016 Oh 2021	0.03; Chi = 3.80 (ree surv 123 123 214 122 171	₽ = 7.00 (P = 0.00) ival 260 198 354 175 288	96 94 184 93 153	243 198 354 215 288	4); I ² = 43% 20.2% 18.5% 22.3% 17.8% 21.1%	0.73 (0.51, 1.04) 0.55 (0.37, 0.82) 0.71 (0.53, 0.95) 0.33 (0.22, 0.50)	
	Heterogeneity: Tau ² = 0 Test for overall effect: Z 1.6.3 5-year disease-fr Bian 2016 Huang 2017 Lin 2021 Lv 2016 Oh 2021 Subtotal (95% CI)	0.03; Chi = 3.80 (ree surv 123 123 214 122 171	[₽] = 7.00 (P = 0.00 198 354 175 288 1275	96 94 94 184 93 153	243 198 354 215 288 1298	4); I [≠] = 43% 20.2% 18.5% 22.3% 17.8% 21.1% 100.0 %	0.73 [0.51, 1.04] 0.55 [0.37, 0.82] 0.71 [0.53, 0.95] 0.33 [0.22, 0.50] 0.78 [0.56, 1.08] 0.61 [0.46, 0.80]	
	Heterogeneity: Tau* = 0 Test for overall effect. Z 1.6.3 5-year disease-fr Bian 2016 Huang 2017 Lin 2021 Lv 2016 Oh 2021 Subtotal (95% CI) Total events	0.03; Chi = 3.80 (123 123 214 122 171 753	P = 7.00 P = 0.01 ival 260 198 354 175 288 1275	96 94 184 93 153 620	P = 0.1 243 198 354 215 288 1298	4); ² = 43% 20.2% 18.5% 22.3% 17.8% 21.1% 100.0%	0.73 [0.51, 1.04] 0.55 [0.37, 0.82] 0.71 [0.53, 0.95] 0.33 [0.22, 0.50] 0.78 [0.56, 1.08] 0.61 [0.46, 0.80]	
	Heterogeneity: Tau* = 0 Test for overall effect. Z 1.6.3 5-year disease-fr Bian 2016 Huang 2017 Lin 2021 Lv 2016 Oh 2021 Subtotal (95% CI) Total events Heterogeneity: Tau* = 0 Test for overall effect?	0.03; Chi = 3.80 (123 123 214 122 171 753 0.07; Chi = 3.54 (P = 7.00 P = 0.00 ival 260 198 354 175 288 1275 P = 12.1 P = 0.00	I, df = 4 (001) 96 94 184 93 153 620 6, df = 4 004)	P = 0.1 243 198 354 215 288 1298 (P = 0.	4); ² = 43% 20.2% 18.5% 22.3% 17.8% 21.1% 100.0% 02); ² = 67 ⁴	0.73 [0.51, 1.04] 0.55 [0.37, 0.82] 0.71 [0.53, 0.95] 0.33 [0.22, 0.50] 0.78 [0.56, 1.08] 0.61 [0.46, 0.80]	+ + + +
	Heterogeneity: Tau" = 0 Test for overall effect: Z 1.6.3 5-year disease-fr Bian 2016 Huang 2017 Lin 2021 Lv 2016 Oh 2021 Subtotal (95% CI) Total events Heterogeneity: Tau" = 0 Test for overall effect: Z	0.03; Chi = 3.80 (123 123 214 122 171 753 0.07; Chi = 3.54 (² = 7.00 (P = 0.01 (P = 0.01 198 354 175 288 1275 ² = 12.1 (P = 0.01 	I, df = 4 (001) 96 94 184 93 153 620 6, df = 4 004)	P = 0.1 243 198 354 215 288 1298 (P = 0.	4); ² = 43% 20.2% 18.5% 22.3% 17.8% 21.1% 100.0% 02); ² = 674	0,73 (0,51, 1,04) 0,55 (0,37, 0,82) 0,71 (0,53, 0,95) 0,33 (0,22, 0,50) 0,78 (0,56, 1,08) 0,61 (0,46, 0,80) %	
	Heterogeneity: Tau ² = 0 Test for overall effect. Z 1.6.3 5-year disease-fr Bian 2016 Huang 2017 Lin 2021 Lv 2016 Oh 2021 Subtotal (95% CI) Total events Heterogeneity: Tau ² = 0 Test for overall effect. Z	0.03; Chi = 3.80 (ree surv 123 123 214 122 171 753 0.07; Chi = 3.54 (P = 7.00 P = 0.00 P = 0.00 198 354 175 288 1275 P = 12.1 P = 0.00	I, df = 4 (001) 96 94 184 93 153 620 6, df = 4 004)	P = 0.1 243 198 354 215 288 1298 (P = 0.	20.2% 18.5% 22.3% 17.8% 21.1% 100.0% 02); I ² = 67 ⁴	0.73 [0.51, 1.04] 0.55 [0.37, 0.82] 0.71 [0.53, 0.95] 0.33 [0.22, 0.50] 0.78 [0.56, 1.08] 0.61 [0.46, 0.80]	

Figure 5 Meta-analysis of DFS in trials comparing No. 10 LND versus non-No. 10 LND for: (A) patients with gastric cancer who have undergone total gastrectomy; (B) patients with gastric cancer and/or Siewert type III AEG who have undergone total gastrectomy. DFS, disease-free survival; No. 10 LND, No. 10 lymph node dissection; AEG, adenocarcinoma of the esophagogastric junction; No. 10+, No. 10 lymph node dissection group; M-H, Mantel-Haenszel; CI, confidence interval.

may be present. Thus, surgeons have preferred splenectomy to facilitate the No. 10 LND for the total gastrectomy in previous years (36,37). However, the total gastrectomy combined with the resection of spleen is reported to result

in higher morbidity, larger blood loss, and could not show a superiority on survival rates compared with that of splenic preservation (38,39). Moreover, the loss of the antitumor and the anti-infection functions of the spleen's immunologic

	No.	10+	No.	10-		Odds ratio (Non-e	event) Ode	ls ratio (Non-event)
Study or subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95%	CI M	-H, Fixed, 95% CI
1.7.2 Complications w	vith Grad	e I-11						
Huang 2017	21	198	15	198	10.1%	1.45 [0.72, 2.90]		
Lin 2021	53	354	46	354	29.3%	1.18 [0.77, 1.80]		
Liu 2021	27	237	23	237	15.3%	1.20 [0.66, 2.15]		
Lv 2016	35	293	29	401	16.2%	1.74 [1.04, 2.92]		
Oh 2021	39	288	45	288	29.2%	0.85 [0.53, 1.34]		
Subtotal (95% CI)		1370		1478	100.0%	1.20 [0.96, 1.51]		◆
Total events	175		158					
Heterogeneity: Chi ² =	4.46, df =	4 (P =	0.35); I ² =	10%				
Test for overall effect:	Z = 1.57 (P = 0.1	2)					
1.7.3 Complications w	ith Grad	e III-IV						
Huang 2017	4	198	5	198	9.2%	0.80 [0.21, 3.01]		
Lin 2021	11	354	7	354	12.8%	1.59 [0.61, 4.15]		
Liu 2021	17	237	14	237	24.5%	1.23 [0.59, 2.56]		
Lv 2016	19	293	11	401	16.4%	2.46 [1.15, 5.25]		
Oh 2021	18	288	21	288	37.1%	0.85 [0.44, 1.63]		
Subtotal (95% CI)		1370		1478	100.0%	1.30 [0.91, 1.85]		•
Total events	69		58					
Heterogeneity: Chi ² = 1	5.08, df =	4 (P =	0.28); I ² =	21%				
Test for overall effect:	Z=1.42 (P = 0.1	6)					
1.7.4 Mortality								
Bian 2016	3	260	1	243	20.5%	2.82 [0.29, 27.34]	6	
Huang 2017	0	198	0	198		Not estimable		~
Lin 2021	0	354	1	354	30.1%	0.33 [0.01, 8.19]		
Liu 2021	3	237	2	237	39.7%	1.51 [0.25, 9.10]	-	
Yang 2014	1	220	0	233	9.7%	3.19 [0.13, 78.76]		
Subtotal (95% CI)		1269		1265	100.0%	1.59 [0.52, 4.87]		
Total events	7		4					
Heterogeneity: Chi ² =	1.35, df =	3 (P =	0.72); l ² =	:0%				
Test for overall effect:	Z = 0.81 (P = 0.4	2)					
							0.02 0.1	
							U.UZ U.I	
							Favours No	10+ Favours No. 10-

Figure 6 Meta-analysis of safety in trials comparing No. 10 LND versus non-No. 10 LND for patients with gastric cancer and/or Siewert type II/III AEG who have undergone total gastrectomy. No. 10 LND, No. 10 lymph node dissection; AEG, adenocarcinoma of the esophagogastric junction; No. 10+, No. 10 lymph node dissection group; No. 10-, non-No. 10 lymph node dissection group; M-H, Mantel-Haenszel; CI, confidence interval.

effect is a negative effect for patients with splenectomy. Sano et al. (40,41) have compared the prognosis of patients with splenectomy and spleen preservation (without intentional No. 10 dissection); found that the 5-year overall survival rates of two arms are 75.1% and 76.4%, respectively (P>0.05); and confirmed the noninferiority of the spleen preservation. However, this study was limited to patients with upper gastric cancer without invasion to the greater curvature. Regarding proximal gastric cancer invading the greater curvature, one retrospective study in Japan reported that no significant survival benefit was observed in the splenectomy group comparing with spleen preservation (without intentional No. 10 dissection) group [5-year OS rate of 63.7% vs. 73.6% and 5-year relapse-free survival (RFS) rate of 60.2% vs. 67.3%], and splenectomy was associated with a higher morbidity rate (30.2% vs. 13.3%) (42). Yang et al. (43) have conducted a metaanalysis and found that splenectomy did not increase 5-year overall survival rate but had significantly higher incidence of postoperative complications. For the above reasons, the laparoscopic spleen-preserving No. 10 LND is first

reported by Huang *et al.* and has been gradually accepted and adopted by an increasing number of surgeons as a technically safe and feasible procedure (44).

In this study, the significant benefit of the No. 10 LND is observed, which is similar to the result reported by Huang et al. (32). Several reasons can account for the decreased mortality. First, the No. 10 LNM is closely linked with the prognosis of patients (15,45). Shin et al. (15) have reported that the 5-year survival rate for patients in the hilar node metastasis group (11.04%) is significantly lower than that in the non-metastasis group (51.57%, P<0.001). Takayama et al. (45) have also reported that the prognosis of patients with positive No. 10 LNs is significantly worse than that of patients with negative No. 10 LNs. Although the No. 10 LND with splenectomy is demonstrated to have no superiority in terms of safety and prognosis over the non-No. 10 LND, many studies have verified that the spleenpreserving No. 10 LND, whether in the laparoscopic or the open form, is safe (46,47). Yang et al. (29) have reported that no spleen-preserving LND-related complication, such as intraperitoneal hemorrhage or pancreatic leakage, has

occurred in the two groups, except one patient from the 10D+ group who has experienced intraoperative splenic injury. Zheng et al. (26) have conducted a prospective multicenter study to evaluate the technical safety and feasibility of laparoscopic spleen-preserving No. 10 LND for 242 patients with total gastrectomy. Results show that the major complication rate is 3.3% (8/242), but No. 10 LND-related complications are not observed (26). Thus, without morbidity and mortality, the spleen-preserving No. 10 LND may bring increased dissection of the positive No. 10 LNs and possibly favorable prognosis. Zheng et al. (26) have reported that the average numbers of laparoscopic spleen-preserving No. 10 LND and metastases are 2.4 and 0.1, respectively, and the rate of the No. 10 LNM is 8.1% among patients with advanced gastric cancer (18/223). Second, the No. 10 LNM is found to be significantly associated with positive No. 4s LN in several studies (30,48,49). Bian et al. (30) have found that the negative predictive efficacy of No. 4s LN status for no metastasis to No. 10 LN is 98.09%. Aoyagi et al. (48) have reported that Nos. 4sa and 4sb LNM are significant parameters for the No. 10 LNM (P<0.001 and P=0.006, respectively) with a logistic regression analysis. Besides, No. 4s LNs are found be upstream of No. 10 LNs (50). Thus, based on the above, the status of negative No. 4s LN may be an indicator for predicting no metastasis to No. 10 LN, and No. 10 LND may gain survival benefits for the patients with No. 4s metastasis. If No. 4s LNs are identified as positive by using intraoperative visualization or pathological examination, the No. 10 LND may be recommended. Third, the greater curvature is found as the common tumor location in patients with No. 10 LNM (38.5%) followed by posterior wall (27.8%), and encircling involvement (22.8%) (4). The serosa-negative tumors located at the lesser curvature and anterior wall are observed with no No. 10 LNM (4). Watanabe et al. (51) have found that the incidence of the No. 10 LNM is 15.9% in patients with tumors invading the greater curvature from a retrospective data of 421 patients' outcomes after the total gastrectomy for proximal advanced gastric cancer, and the index to estimate the benefit from the No. 10 LND is 5.6, indicating a certain survival benefit. The greater curvature invasion is verified to be a risk factor of the No. 10 LNM, and therefore it had clinical significance of No. 10 LND for these patients with advanced gastric cancer invading the greater curvature line (38,51). Maezawa et al. (52) have found the substantially higher T1 of the No. 10 LN than the other regional nodes, such as Nos. 8a, 11p, and 11d, in locally advanced proximal

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gastric cancer invading the greater curvature. Authors recommend the No. 10 LND as a part of the D2 dissection for proximal gastric cancer invading the greater curvature (52).

The meta-analyses show that the DFS rates for patients with the No. 10 LND have significantly better prognosis compared with those for patients without the No. 10 LND. Three studies are not included in the meta-analyses because of insufficient data (14,29,34). However, Park et al. (14) have observed a RFS rate and found no difference in the RFS rates between the two groups. The DFS rate, which is closely correlated to relapse and distant metastasis, generally denotes the length of time the patient survives without any signs or symptoms after primary treatment for a cancer (53). The presence of the No. 10 LNM is one of the independent predictors of distant metastasis after the R0 surgical resection, indicating that the non-No. 10 LND patients with potentially positive No. 10 LN have a high risk for the presence of distant LNM and therefore a lower DFS, but a limited risk for the presence of relapse in situ (6). Thus, the relative effect of the No. 10 LND may be less remarkable for RFS than DFS.

Considering the discrepancies of Siewert type II AEG, Siewert type III AEG, and stomach cancer, subgroup analysis according to the location of tumor are performed. The results of subset analyses showed that the survival and DFS rates of the patients with gastric cancer are consistent with those of the patients with gastric cancer and/or type II/ III AEG. Results indicate that the tumor, whether located in the stomach or below 1 centimeter above the esophagogastric junction, is not crucial in the prognosis of patients. The finding maybe because the biological behavior and the anatomical position of the Siewert type II/III AEG are quite similar to those of advanced proximal gastric cancer, which has a similar prognosis while the total gastrectomy and the D2 lymphadenectomy are performed (54). The subgroup analyses on the tumor of gastric cancer and/or Siewert type III AEG indicate that the No. 10 LND significantly improves the overall survival and the DFS rates of patients with gastric cancer and/or the Siewert type III AEG who have undergone total gastrectomy. However, the validity of meta-analysis may be affected by significant heterogeneity and limited sample sizes. The subset analyses according to tumor size, tumor stage, and degree of pathological differentiation are not conducted because of insufficient information. The result of meta-analysis of safety in trials comparing No. 10 LND versus non-No. 10 LND indicates that No. 10 LND is a safe way for patients with gastric cancer and/or Siewert type II/III AEG who have undergone

total gastrectomy.

The current study has several potential limitations. First, there are important heterogeneities among studies. Sensitivity and subgroup analyses have failed to eliminate the significance. There are many differences across studies that serve as sources of heterogeneity, including variation in sample sizes, variation in the baseline of tumor characteristics (e.g., tumor differentiation, stage, and size), and length of follow-up period. Second, the publication bias may exist because of the relatively limited database. The quality of the current study may be influenced by none of the available RCTs included. We speculate that the reason may be that for surgeons, performing surgical intervention with randomized and blinded ways is a difficult and unethical task. Nevertheless, the meta-analyses of well-designed non-RCTs are demonstrated to have similar accuracy to RCTs (55). There may be duplicate patients due to several overlaps in terms of institution and operation year, in which the patients underwent total gastrectomy (29,31-34). However, we failed to eliminate the duplicate data because of the insufficient reported information in the included studies. Nevertheless, we conducted sensitivity analyses of the studies that may have contained duplicate patients to verify the stability of the results. The analysis results indicated that the survival outcomes in this study are slightly influenced by potential duplicate patients. Besides, there may exist potentially confounding selection bias from the non-No. 10 LND group introduced by the retrospective nature of the included studies, because it is difficult to analyze and draw a conclusion regarding the No. 10 LN metastasis and staging without lymphadenectomy. The improved comprehensive approach of imaging diagnosis, intraoperative diagnosis on LN metastasis and staging can be expected in the future (56). In addition, the chemotherapy is a prognostic factor of patients with gastric cancer and/or Siewert type II/III AEG who have undergone the total gastrectomy. Hence, patients who have received postoperative chemotherapy during the period of the clinical researches may influence the results. Therefore, the results of current study should be interpreted cautiously.

Conclusions

The No. 10 LND with spleen preservation is a safe approach to improve the survival of patients with gastric cancer and/or Siewert type II/III AEG who have undergone total gastrectomy. Further high-quality prospective trials are urgently needed to verify this outcome.

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Footnote

Reporting Checklist: The authors have completed the PRISMA reporting checklist. Available at https://tcr. amegroups.com/article/view/10.21037/tcr-22-522/rc

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Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at https://tcr.amegroups.com/article/view/10.21037/tcr-22-522/coif). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

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References

- Joshi SS, Badgwell BD. Current treatment and recent progress in gastric cancer. CA Cancer J Clin 2021;71:264-79.
- Sung H, Ferlay J, Siegel RL, et al. Global Cancer Statistics 2020: GLOBOCAN Estimates of Incidence and Mortality Worldwide for 36 Cancers in 185 Countries. CA Cancer J Clin 2021;71:209-49.
- Japanese Gastric Cancer Association. Japanese gastric cancer treatment guidelines 2010 (ver. 3). Gastric Cancer 2011;14:113-23.
- 4. Sasada S, Ninomiya M, Nishizaki M, et al. Frequency of

Xia et al. No. 10 LND for total gastrectomy: a systematic review

lymph node metastasis to the splenic hilus and effect of splenectomy in proximal gastric cancer. Anticancer Res 2009;29:3347-51.

- Nashimoto A, Yabusaki H, Matsuki A. The significance of splenectomy for advanced proximal gastric cancer. Int J Surg Oncol 2012;2012:301530.
- 6. Zhu GL, Sun Z, Wang ZN, et al. Splenic hilar lymph node metastasis independently predicts poor survival for patients with gastric cancers in the upper and/or the middle third of the stomach. J Surg Oncol 2012;105:786-92.
- Guner A, Hyung WJ. Advantages of Splenic Hilar Lymph Node Dissection in Proximal Gastric Cancer Surgery. J Gastric Cancer 2020;20:19-28.
- Kakeji Y, Yamamoto M, Ito S, et al. Lymph node metastasis from cancer of the esophagogastric junction, and determination of the appropriate nodal dissection. Surg Today 2012;42:351-8.
- Goto H, Tokunaga M, Sugisawa N, et al. Value of splenectomy in patients with Siewert type II adenocarcinoma of the esophagogastric junction. Gastric Cancer 2013;16:590-5.
- Hasegawa S, Yoshikawa T, Rino Y, et al. Priority of lymph node dissection for Siewert type II/III adenocarcinoma of the esophagogastric junction. Ann Surg Oncol 2013;20:4252-9.
- Kurokawa Y, Takeuchi H, Doki Y, et al. Mapping of Lymph Node Metastasis From Esophagogastric Junction Tumors: A Prospective Nationwide Multicenter Study. Ann Surg 2021;274:120-7.
- Japanese Gastric Cancer Association. Japanese classification of gastric carcinoma: 3rd English edition. Gastric Cancer 2011;14:101-12.
- Galizia G, Lieto E, De Vita F, et al. Modified versus standard D2 lymphadenectomy in total gastrectomy for nonjunctional gastric carcinoma with lymph node metastasis. Surgery 2015;157:285-96.
- Park SH, Son T, Seo WJ, et al. Prognostic Impact of Extended Lymph Node Dissection versus Limited Lymph Node Dissection on pN0 Proximal Advanced Gastric Cancer: a Propensity Score Matching Analysis. J Gastric Cancer 2019;19:212-24.
- Shin SH, Jung H, Choi SH, et al. Clinical significance of splenic hilar lymph node metastasis in proximal gastric cancer. Ann Surg Oncol 2009;16:1304-9.
- Japanese Gastric Cancer Association. Japanese gastric cancer treatment guidelines 2018 (5th edition). Gastric Cancer 2021;24:1-21.
- 17. Page MJ, McKenzie JE, Bossuyt PM, et al. The PRISMA

2020 statement: An updated guideline for reporting systematic reviews. J Clin Epidemiol 2021;134:178-89.

- Wells GA, Shea B, O'Connell D, et al. The Newcastle-Ottawa Scale (NOS) for assessing the quality of nonrandomised studies in meta-analyses. Available online: http://www.ohri.ca/programs/clinical_epidemiology/ oxford.htm
- 19. Jadad AR, Moore RA, Carroll D, et al. Assessing the quality of reports of randomized clinical trials: is blinding necessary? Control Clin Trials 1996;17:1-12.
- Egger M, Davey Smith G, Schneider M, et al. Bias in meta-analysis detected by a simple, graphical test. BMJ 1997;315:629-34.
- 21. Begg CB, Mazumdar M. Operating characteristics of a rank correlation test for publication bias. Biometrics 1994;50:1088-101.
- 22. Lin JX, Huang CM, Zheng CH, et al. Is it necessary to dissect the posterior lymph nodes along the splenic vessels during total gastrectomy with D2 lymphadenectomy for advanced gastric cancer? Eur J Surg Oncol 2017;43:2357-65.
- 23. Yang K, Lu ZH, Zhang WH, et al. Comparisons Between Different Procedures of No. 10 Lymphadenectomy for Gastric Cancer Patients With Total Gastrectomy. Medicine (Baltimore) 2015;94:e1305.
- Ji X, Fu T, Bu ZD, et al. Comparison of different methods of splenic hilar lymph node dissection for advanced upper- and/or middle-third gastric cancer. BMC Cancer 2016;16:765.
- 25. Wang JB, Liu ZY, Chen QY, et al. Short-term efficacy of robotic and laparoscopic spleen-preserving splenic hilar lymphadenectomy via Huang's three-step maneuver for advanced upper gastric cancer: Results from a propensity score-matched study. World J Gastroenterol 2019;25:5641-54.
- Zheng CH, Xu YC, Zhao G, et al. Safety and feasibility of laparoscopic spleen-preserving No. 10 lymph node dissection for locally advanced upper third gastric cancer: a prospective, multicenter clinical trial. Surg Endosc 2020;34:5062-73.
- 27. Zheng C, Xu Y, Zhao G, et al. Outcomes of Laparoscopic Total Gastrectomy Combined With Spleen-Preserving Hilar Lymphadenectomy for Locally Advanced Proximal Gastric Cancer: A Nonrandomized Clinical Trial. JAMA Netw Open 2021;4:e2139992.
- Zhong Q, Chen QY, Xu YC, et al. Reappraise role of No. 10 lymphadenectomy for proximal gastric cancer in the era of minimal invasive surgery during total gastrectomy: a pooled analysis of 4 prospective trial. Gastric Cancer

3036

2021;24:245-57.

- Yang K, Zhang WH, Chen XZ, et al. Survival benefit and safety of no. 10 lymphadenectomy for gastric cancer patients with total gastrectomy. Medicine (Baltimore) 2014;93:e158.
- 30. Bian S, Xi H, Wu X, et al. The Role of No. 10 Lymphadenectomy for Advanced Proximal Gastric Cancer Patients Without Metastasis to No. 4sa and No. 4sb Lymph Nodes. J Gastrointest Surg 2016;20:1295-304.
- 31. Lv CB, Huang CM, Zheng CH, et al. Should Splenic Hilar Lymph Nodes be Dissected for Siewert Type II and III Esophagogastric Junction Carcinoma Based on Tumor Diameter?: A Retrospective Database Analysis. Medicine (Baltimore) 2016;95:e3473.
- 32. Huang CM, Chen T, Lin JX, et al. The effects of laparoscopic spleen-preserving splenic hilar lymphadenectomy on the surgical outcome of proximal gastric cancer: a propensity score-matched, case-control study. Surg Endosc 2017;31:1383-92.
- Lin JX, Wang ZK, Huang YQ, et al. Clinical Relevance of Splenic Hilar Lymph Node Dissection for Proximal Gastric Cancer: A Propensity Score-Matching Case-Control Study. Ann Surg Oncol 2021;28:6649-62.
- Liu K, Chen XZ, Zhang YC, et al. The value of spleenpreserving lymphadenectomy in total gastrectomy for gastric and esophagogastric junctional adenocarcinomas: A long-term retrospective propensity score match study from a high-volume institution in China. Surgery 2021;169:426-35.
- 35. Oh YJ, Kim DH, Eom BW, et al. Is Splenic Hilar Lymph Node Dissection Without Splenectomy Essential for Proximal Advanced Gastric Cancer? Ann Surg Oncol 2021;28:8952-61.
- Nakata K, Nagai E, Ohuchida K, et al. Technical feasibility of laparoscopic total gastrectomy with splenectomy for gastric cancer: clinical short-term and long-term outcomes. Surg Endosc 2015;29:1817-22.
- Csendes A, Burdiles P, Rojas J, et al. A prospective randomized study comparing D2 total gastrectomy versus D2 total gastrectomy plus splenectomy in 187 patients with gastric carcinoma. Surgery 2002;131:401-7.
- Toriumi T, Terashima M. Disadvantages of Complete No. 10 Lymph Node Dissection in Gastric Cancer and the Possibility of Spleen-Preserving Dissection: Review. J Gastric Cancer 2020;20:1-18.
- Kinoshita T, Okayama T. Is splenic hilar lymph node dissection necessary for proximal gastric cancer surgery? Ann Gastroenterol Surg 2020;5:173-82.

- Sano T, Sasako M, Mizusawa J, et al. Randomized Controlled Trial to Evaluate Splenectomy in Total Gastrectomy for Proximal Gastric Carcinoma. Ann Surg 2017;265:277-83.
- Sano T, Yamamoto S, Sasako M, et al. Randomized controlled trial to evaluate splenectomy in total gastrectomy for proximal gastric carcinoma: Japan clinical oncology group study JCOG 0110-MF. Jpn J Clin Oncol 2002;32:363-4.
- 42. Ohkura Y, Haruta S, Shindoh J, et al. Efficacy of prophylactic splenectomy for proximal advanced gastric cancer invading greater curvature. World J Surg Oncol 2017;15:106.
- Yang K, Zang ZY, Niu KF, et al. The Survival Benefit and Safety of Splenectomy for Gastric Cancer With Total Gastrectomy: Updated Results. Front Oncol 2021;10:568872.
- 44. Huang CM, Chen QY, Lin JX, et al. Huang's three-step maneuver for laparoscopic spleen-preserving No. 10 lymph node dissection for advanced proximal gastric cancer. Chin J Cancer Res 2014;26:208-10.
- 45. Takayama T, Wakatsuki K, Matsumoto S, et al. Prognostic significance of splenic hilar nodal involvement in proximal third gastric carcinoma. Hepatogastroenterology 2011;58:647-51.
- Oh SJ, Hyung WJ, Li C, et al. The effect of spleenpreserving lymphadenectomy on surgical outcomes of locally advanced proximal gastric cancer. J Surg Oncol 2009;99:275-80.
- 47. Mou TY, Hu YF, Yu J, et al. Laparoscopic splenic hilum lymph node dissection for advanced proximal gastric cancer: a modified approach for pancreas- and spleenpreserving total gastrectomy. World J Gastroenterol 2013;19:4992-9.
- Aoyagi K, Kouhuji K, Miyagi M, et al. Prognosis of metastatic splenic hilum lymph node in patients with gastric cancer after total gastrectomy and splenectomy. World J Hepatol 2010;2:81-6.
- 49. Ikeguchi M, Kaibara N. Lymph node metastasis at the splenic hilum in proximal gastric cancer. Am Surg 2004;70:645-8.
- 50. Takahashi T, Sawai K, Hagiwara A, et al. Type-oriented therapy for gastric cancer effective for lymph node metastasis: management of lymph node metastasis using activated carbon particles adsorbing an anticancer agent. Semin Surg Oncol 1991;7:378-83.
- Watanabe M, Kinoshita T, Enomoto N, et al. Clinical Significance of Splenic Hilar Dissection with Splenectomy

Xia et al. No. 10 LND for total gastrectomy: a systematic review

in Advanced Proximal Gastric Cancer: An Analysis at a Single Institution in Japan. World J Surg 2016;40:1165-71.

- 52. Maezawa Y, Aoyama T, Yamada T, et al. Priority of lymph node dissection for proximal gastric cancer invading the greater curvature. Gastric Cancer 2018;21:569-72.
- 53. National Cancer Institute. NCI Dictionary of Cancer Terms. Available online: https://www.cancer.gov/ publications/dictionaries/cancer-terms/def/disease-freesurvival
- 54. Mönig SP, Collet PH, Baldus SE, et al. Splenectomy in

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proximal gastric cancer: frequency of lymph node metastasis to the splenic hilus. J Surg Oncol 2001;76:89-92.

- 55. Abraham NS, Byrne CJ, Young JM, et al. Meta-analysis of well-designed nonrandomized comparative studies of surgical procedures is as good as randomized controlled trials. J Clin Epidemiol 2010;63:238-45.
- 56. Lee S, Song JH, Choi S, et al. Fluorescent lymphography during minimally invasive total gastrectomy for gastric cancer: an effective technique for splenic hilar lymph node dissection. Surg Endosc 2022;36:2914-24.

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