Hindawi BioMed Research International Volume 2020, Article ID 6131968, 10 pages https://doi.org/10.1155/2020/6131968

# Review Article

# Roux-en-Y and Billroth II Reconstruction after Pancreaticoduodenectomy: A Meta-Analysis of Complications

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Received 14 August 2020; Revised 9 November 2020; Accepted 18 November 2020; Published 4 December 2020

Academic Editor: Celal Ulasoglu

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Objective. To evaluate Roux-en-Y and Billroth II reconstruction following pancreaticoduodenectomy (PD). Methods. PubMed, Embase, the Cochrane Library, and the Web of Science were searched to identify randomized controlled trials (RCTs) and controlled clinical trials that compared Roux-en-Y and Billroth II reconstruction following PD up to December 2019. RevMan 5.3 software was used for the statistical analysis. Results. Four RCTs and five controlled clinical trials were included, with a total of 1,072 patients (500 and 572 patients in the Roux-en-Y and Billroth II groups, respectively). No significant differences in delayed gastric emptying (DGE), A-grade DGE, B-grade DGE, or C-grade DGE were observed between the Roux-en-Y and Billroth II reconstruction groups after PD (odds ratio [OR] = 1.01, 95% confidence interval [CI]: 0.50–2.03, P = 0.98; OR = 0.49, 95% CI: 0.17–1.45, P = 0.20; OR = 0.63, 95% CI: 0.29–1.38, P = 0.25; and OR = 2.13, 95% CI: 0.38–11.99, P = 0.39). No significant difference in the incidence of postoperative pancreatic fistula, abscess, bile leaks, infection, postoperative bleeding, or the length of the postoperative hospital stay was observed between the Roux-en-Y and Billroth II groups (P > 0.05), but the operation time was significantly different (mean difference [MD] = 31.65, 95% CI: 7.14–56.17, P = 0.01). Conclusions. Billroth II reconstruction after PD did not significantly reduce the incidence of DGE or other complications but shortened the operation time compared to Roux-en-Y reconstruction. However, the results must be verified by further high-quality, large RCTs or controlled clinical trials.

#### 1. Introduction

The incidence of pancreatic cancer is rising, and it is estimated that pancreatic cancer will become the second most deadly cancer in the world by 2030 [1]. Scholars have performed a considerable amount of research on pancreatico-duodenectomy (PD) since Whipple et al. first proposed the concept in 1935 [2]. Complications after PD have a significant impact on the postoperative quality of life of patients, for whom the total and clinically relevant delayed gastric emptying (DGE) incidence rates are 27.7% and 14.3%, respectively. The incidence of DGE is associated with the pancreatic resection type, pylorus preservation status (yes

or no), antecolic and retrocolic gastrojejunal anastomosis, and gastrojejunal anastomosis type [3]. The incidence of DGE after Billroth II and Roux-en-Y gastrojejunal reconstruction has remained controversial. Based on the hypothesis that Roux-en-Y anatomy can prevent gastric contents from activating trypsin, Machado et al. [4] proposed that Roux-en-Y "protects" the pancreatojejunostomy. Results of single-center randomized controlled trials (RCTs) comparing the incidence of DGE between Roux-en-Y and Billroth II reconstruction are not consistent [5–8]. Meta-analyses have also reported contrasting results. Yang et al. [9] proposed that Billroth II reconstruction reduces the incidence of clinical DGE compared to Roux-en-Y; however, Li et al.

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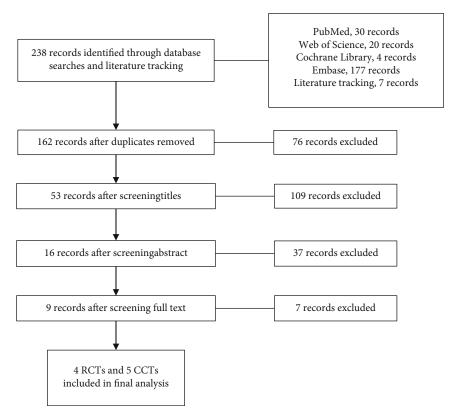


FIGURE 1: PRISMA flow chart of the literature screening process.

Table 1: Basic parameters of the included studies.

Study	Publication time	Number in study	R/B (n)	Age (years) Roux-en-Y/Billroth II	Gender (M/F)	Study period	Study type	Nation
Shimoda et al. [5]	2013	101	49/52	65.7 ± 11.1/66.5 ± 9.8	60/41	2008-2011	RCT	Japan
Busquets et al. [6]	2018	80	40/40	$68.1 \pm 11.7$ $/65.6 \pm 10.9$	48/32	2013-2015	RCT	Spain
Tani et al. [7]	2014	151	75/76	$69.6 \pm 7.9/68.0 \pm 8.9$	81/70	2009-2012	RCT	Japan
Ke et al. [8]	2013	216	107/109	$58.3 \pm 5.9 / 59.3 \pm 6.6$	101/115	2006-2012	RCT	China
Perwaiz et al. [13]	2009	108	53/55	$53.3 \pm 12.1$ $/53.5 \pm 10.1$	81/27	2003-2007	CCT	India
Kaman et al. [14]	2008	111	60/51	$51 \pm 13.3/50 \pm 13.6$	74/37	1994-2006	CCT	India
Ballas et al. [15]	2010	88	46/42	$64.4 \pm 9.5/60.9 \pm 11.5$	52/36	1994-2006	CCT	Greece
Ben-Ishay et al. [16]	2019	179	52/127	$68.2 \pm 9.6/68 \pm 13.7$	88/91	2010-2016	CCT	Israel
Casadei et al. [17]	2008	38	18/20	$65.7 \pm 10.0$ $/56.3 \pm 11.0$	24/14	2006-2007	CCT	Italy

R: Roux-en-Y; B: Billroth II; M: male; F: female.

[10] reported no difference between the two reconstruction types.

The purpose of this study is to evaluate the ability of Roux-en-Y and Billroth II reconstruction following PD to prevent DGE and other complications.

## 2. Materials and Methods

#### 2.1. Selection Criteria

2.1.1. Inclusion Criteria. The inclusion criteria are as follows: (1) a clinical comparative study between Roux-en-Y and Billroth II gastrojejunal reconstruction following PD, (2) RCT or clinical controlled trial (CCT), and (3) English.

2.1.2. Exclusion Criteria. The exclusion criteria are as follows: (1) no outcome indicators, (2) pancreaticogastrostomy, (3) reviews and case reports, and (4) gastrectomy history.

Table 2: Pathologies, surgery-related parameters, and quality scores.

Study		Patho	logies (	R/B)		PJ & GJ	Stenting	Definition of DGE	Definition	QS
	PL	ВС	AC	DC	Other		(yes/no)		of POPF	-Q0
Shimoda et al. [5]						Duct-to-mucosa, end-to-side	Yes	ISGPS2007	ISGPF2005	5
Busquets et al. [6]	28/28	5/3	6/9	1/0		Duct-to-mucosa, end-to-side		ISGPS2007	ISGPF2005	5
Tani et al. [7]	55/57	11/11	7/6	2/2		Duct-to-mucosa, end-to-side	Yes	ISGPS2007	ISGPF2005	5
Ke et al. [8]	51/50	32/35	18/16	6/8		Duct-to-mucosa, end-to-side	Yes	Johns Hopkins [18]	Johns Hopkins [19], ISGPF2005	5
Perwaiz et al. [13]	12/13	10/4	24/25	3/6	4/7	Duct-to-mucosa, end-to-side	Yes	van Berge Henegouwen [20]	ISGPF2005	8
Kaman et al. [14]	30/24	9/4	16/18	5/5		Mucosa-to- mucosa, end-to-side	Yes	Self-definition	Self-definition	8
Ballas et al. [15]	59*	8*	15*	6*		Duct-to-mucosa, end-to-side	Yes	_	ISGPF2005	6
Ben-Ishay et al. [16]						End-to-side		ISGPS2007	_	8
Casadei et al. [17]	12/20	1/0	5/0			Duct-to-mucosa, end-to-side	Yes	_	ISGPF2005	7

R: Roux-en-Y; B: Billroth II; PL: pancreatic lesions; BC: biliary cancer; AC: ampullary cancer; DC: duodenal cancer; PJ: pancreatojejunostomy; GJ: gastrojejunostomy; QS: quality score; ISGPS: International Study Group of Pancreatic Surgery; ISGPF: International Study Group for Pancreatic Fistula; —: not mentioned. \*total number (R+B).

	Roux-	en-Y	Billrot	h II		Odds ratio		Odds	ratio	
Study or subgroup	Events	Total	Events	Total	Weight	M-H, random, 95%	CI	M-H, rand	lom, 95% CI	
2.1.1 RCT										
Busquets 2018	18	40	18	40	13.1%	1.00 [0.41, 2.41]				
Ke 2013	25	107	27	109	14.6%	0.93 [0.50, 1.73]		-	_	
Shimoda 2013	10	49	3	52	10.2%	4.19 [1.08, 16.27]			-	
Tani 2014	11	75	9	76	12.7%	1.28 [0.50, 3.29]		-	-	
Subtotal (95% CI)		271		277	50.5%	1.23 [0.73, 2.07]		•		
Total events	64		57							
Heterogeneity: tau <sup>2</sup>	= 0.08; c	$hi^2 = 4$	.10, df =	3 (P =	$= 0.25); I^2 =$	27%				
Test for overall effec	t: $Z = 0.7$	78 (P =	0.44)							
2.1.2 CCT										
Ballas 2010	7	46	4	42	10.5%	1.71 [0.46, 6.30]		-		
Ben-Ishay 2019	8	52	75	127	13.4%	0.13 [0.05, 0.29]		<del></del>		
Casadei 2008	0	18	1	20	3.6%	0.35 [0.01, 9.18]		•		
kaman 2008	12	60	6	51	12.0%	1.88 [0.65, 5.42]		-		
perwaiz 2009	5	53	4	55	10.1%	1.33 [0.34, 5.24]		-	-	
Subtotal (95% CI)		229		295	49.5%	0.75 [0.19, 2.91]				
Total events	32		90							
Heterogeneity: tau <sup>2</sup>	= 1.82; c	$hi^2 = 2$	1.83, df	= 4 (P	= 0.0002);	$I^2 = 82\%$				
Test for overall effec										
Total (95% CI)		500		572	100.0%	1.01 [0.50, 2.03]			<b>&gt;</b>	
Total events	96		147							
Heterogeneity: tau <sup>2</sup>	= 0.77; c	$hi^2 = 3$	0.50, df	= 8 (P	= 0.0002);	$I^2 = 74\%$			-	
Test for overall effec				- \-			0.01	0.1 1	10	10
Test for subgroup di		,	,	c 1 /	D 0.50) I	2 004	Favou	rs [Roux-en-Y]	Favours [Billro	th II]

FIGURE 2: Forest plot of the incidence of delayed gastric emptying (DGE).

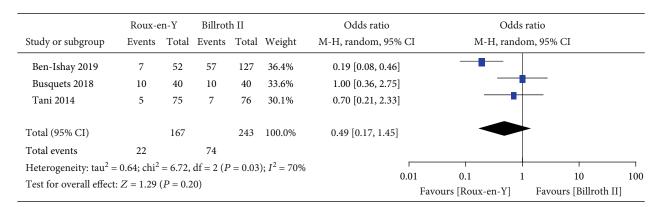


FIGURE 3: Forest plot of the incidence of A-grade delayed gastric emptying (DGE).

	Roux-e	en-Y	Billrot	h II		Odds ratio	(	Odds ra	tio	
Study or subgroup	Events	Total	Events	Total	Weight	M-H, fixed, 95% CI	M-H	I, fixed,	95% CI	
Ben-Ishay 2019	1	52	15	127	51.5%	0.15 [0.02, 1.14]		-		
Busquets 2018	6	40	5	40	25.6%	1.24 [0.34, 4.43]	_			
Shimoda 2013	2	49	2	52	11.2%	1.06 [0.14, 7.86]				
Tani 2014	2	75	2	76	11.7%	1.01 [0.14, 7.39]		1		
Total (95% CI)		216		295	100.0%	0.63 [0.29, 1.38]	<			
Total events	11		24							
Heterogeneity: chi <sup>2</sup> =	= 3.50, df =	3 (P =	0.32); I <sup>2</sup>	= 14%		0.01	0.1	+	10	100
Test for overall effect	z: Z = 1.16	(P = 0.2)	25)			0.01 Fa	0.1 vours [Roux-en-	1 Y]	10 Favours [Billroth	100 II]

FIGURE 4: Forest plot of the incidence of B-grade delayed gastric emptying (DGE).

	Roux-e	en-Y	Billrot	h II		Odds ratio		Odds	ratio	
Study or subgroup	Events	Total	Events	Total	Weight	M-H, random, 95% CI		M-H, rando	om, 95% CI	
Ben-Ishay 2019	0	52	3	127	20.0%	0.34 [0.02, 6.67]		-		
Busquets 2018	2	40	3	40	31.5%	0.65 [0.10, 4.11]				
Shimoda 2013	8	49	1	52	28.3%	9.95 [1.20, 82.83]				
Tani 2014	4	75	0	76	20.3%	9.63 [0.51, 182.05]			•	<b></b>
Total (95% CI)		216		295	100.0%	2.13 [0.38, 11.99]				
Total events	14		7							
Heterogeneity: tau <sup>2</sup> =	1.58; chi <sup>2</sup>	= 6.20,	df = 3 (P	P = 0.10	); $I^2 = 52\%$	б	0.1	0.1	10	100
Test for overall effect:	Z = 0.86	(P = 0.3)	9)			0.0		0.1 rs [Roux-en-Y]	l 10 Favours [Bil	lroth II]

FIGURE 5: Forest plot of the incidence of C-grade delayed gastric emptying (DGE).

2.2. Search Strategy and Screening Methods. PubMed, Embase, the Cochrane Library, and the Web of Science were searched up to December 2019. The search terms were as follows: ((((((pancreatoduodenectomy) OR pancreaticoduodenectomy) OR Whipple))AND(((Roux-en-Y) OR double loop) OR dual loop))AND((((Billroth II) OR conventional reconstruction) OR conventional loop reconstruction) OR single loop))AND(((delayed gastric emptying) OR pancreatic fistula) OR postoperative pancreatic fistula). Published

RCTs and CCTs comparing Roux-en-Y and Billroth II reconstruction following PD were searched. Two researchers independently screened the studies, cross-checked their quality, and asked a third researcher to settle any controversies regarding whether to include a study.

2.3. Quality Assessment. The modified Jadad Scale [11] and Newcastle-Ottawa Scale [12] were used to assess the quality of the RCTs and CCTs, respectively.

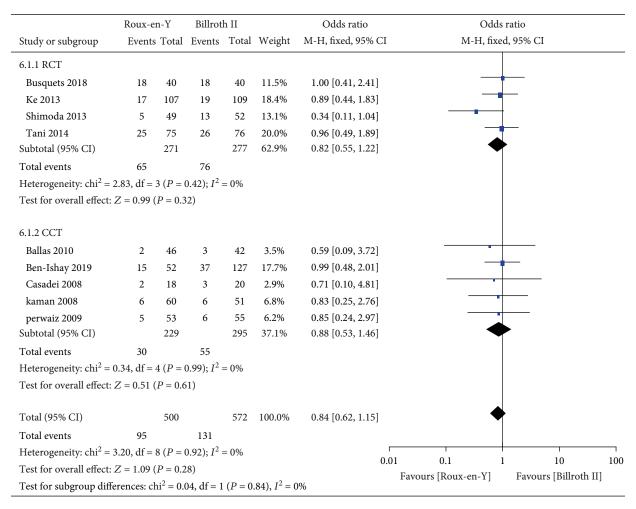


FIGURE 6: Forest plot of the incidence of postoperative pancreatic fistula (POPF).

	Roux-e	en-Y	Billrot	h II		Odds ratio		Od	ds ratio		
Study or subgroup	Events	Total	Events	Total	Weight	M-H, fixed, 95%	CI	M-H, fix	ed, 95% CI		
Ben-Ishay 2019	8	52	14	127	22.2%	1.47 [0.58, 3.74]	]		-		
Busquets 2018	9	40	8	40	20.0%	1.16 [0.40, 3.40]	]	_	-		
kaman 2008	13	60	6	51	16.4%	2.07 [0.73, 5.93]	]		+-	_	
Ke 2013	4	107	4	109	12.3%	1.02 [0.25, 4.19]	]		+		
Tani 2014	7	75	10	76	29.1%	0.68 [0.24, 1.89]	]		•		
Total (95% CI)		334		403	100.0%	1.22 [0.76, 1.96]	]		•		
Total events	41		42								
Heterogeneity: chi <sup>2</sup> =	2.46, df =	4 (P =	0.65); I <sup>2</sup>	= 0%					-		
Test for overall effect	z = 0.83	(P = 0.4)	1)				0.01	0.1	1	10	100
			,				Favoi	ırs [Roux-en-Y]	Favo	urs [Billroth	n II]

FIGURE 7: Forest plot of the incidence of abscess.

2.4. Data Extraction. The general information extracted included the first author's name, year of publication, number of patients included in the study, their age and gender, and the study period, study type, and country. The primary outcomes were DGE, A-grade DGE, B-grade DGE, and C-grade DGE, and the secondary outcomes were post-

operative pancreatic fistula (POPF), abscess, bile leak, infection, postoperative bleeding, operation time, and length of postoperative hospital stay.

2.5. Statistical Analysis. RevMan 5.3 software (Cochrane Collaboration, Copenhagen, Denmark) was used for the statistical

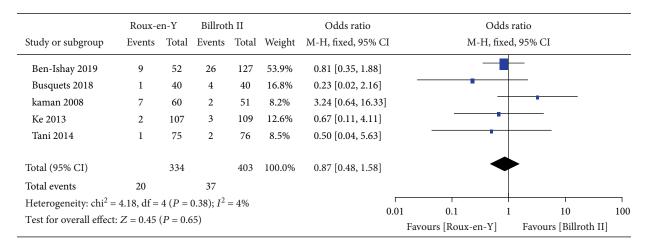


FIGURE 8: Forest plot of the incidence of bile leak.

	Roux-e	en-Y	Billrot	h II		Odds ratio	Odds	ratio	
Study or subgroup	Events	Total	Events	Total	Weight	M-H, fixed, 95% CI	M-H, fixed	, 95% CI	
Ballas 2010	3	46	2	42	3.8%	1.40 [0.22, 8.79]		-	
Ben-Ishay 2019	16	52	28	127	21.9%	1.57 [0.76, 3.24]	+	-	
Casadei 2008	3	18	0	20	0.8%	9.26 [0.44, 192.72]	-	•	<b>→</b>
kaman 2008	17	60	15	51	22.6%	0.95 [0.42, 2.16]		<del></del>	
Ke 2013	19	107	23	109	36.4%	0.81 [0.41, 1.59]	-	_	
perwaiz 2009	5	53	3	55	5.2%	1.81 [0.41, 7.96]		•	
Tani 2014	2	75	5	76	9.4%	0.39 [0.07, 2.07]	•		
Total (95% CI)		411		480	100.0%	1.10 [0.76, 1.61]	•	•	
Total events	65		76						
Heterogeneity: chi <sup>2</sup> =	5.73, df =	6 (P =	$0.45); I^2$	= 0%			1	1	
Test for overall effect	z = 0.52	(P=0.6	50)			0.01	0.1 1	10	100
		•	•			F	avours [Roux-en-Y]	Favours [Billroth	Ш

 $\ensuremath{\mathsf{Figure}}$  9: Forest plot of the incidence of infection.

	Roux-e	en-Y	Billrot	h II		Odds ratio	Odds 1	ratio	
Study or subgroup	Events	Total	Events	Total	Weight	M-H, fixed, 95% CI	M-H, fixed	, 95% CI	
Ballas 2010	1	46	1	42	9.0%	0.91 [0.06, 15.04]	•		
Busquets 2018	3	40	5	40	40.8%	0.57 [0.13, 2.55]			
kaman 2008	7	60	3	51	25.3%	2.11 [0.52, 8.64]	-	_	
perwaiz 2009	2	53	2	55	16.7%	1.04 [0.14, 7.66]			
Tani 2014	4	75	1	76	8.3%	4.23 [0.46, 38.72]		•	
Total (95% CI)		274		264	100.0%	1.37 [0.64, 2.95]	•	<b>&gt;</b>	
Total events	17		12						
Heterogeneity: $chi^2 = 2$	2.83, df =	4 (P = 0	0.59); I <sup>2</sup>	= 0%		0.01	1	10	_
Test for overall effect:	Z = 0.81	(P = 0.4)	12)			0.01 Fa	0.1 1 vours [Roux-en-Y]	10 Favours [Billroth II]	100

Figure 10: Forest plot of the incidence of postoperative bleeding.

	Rou	ıx-en-	Y	Bill	roth I	I		Mean difference	Mean difference
Study or subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, random, 95% CI	IV, random, 95% CI
11.1.1 RCT									
Busquets 2018	431	85	40	405	88	40	10.7%	26.00 [-11.92, 63.92]	-
Ke 2013	342	24	107	336	24	109	14.2%	6.00 [-0.40, 12.40]	-
Shimoda 2013	471	81	49	471	65	52	12.0%	0.00 [-28.75, 28.75]	
Tani 2014	427	131	75	395	68	76	11.4%	32.00 [-1.36, 65.36]	•
Subtotal (95% CI)			271			277	48.3%	8.68 [-1.06, 18.42]	•
Heterogeneity: tau <sup>2</sup> =	22.09; cl	$hi^2 = 3$	.44, df	= 3 (P =	= 0.33)	$; I^2 = 1$	.3%		
Test for overall effect	Z = 1.75	5 (P =	0.08)						
11.1.2 CCT									
Ballas 2010	366.1	60.1	46	333.8	52.7	42	12.7%	32.30 [8.73, 55.87]	<del></del>
Casadei 2008	343	38	18	331	33.5	20	12.8%	12.00 [-10.89, 34.89]	<del>    •</del>
kaman 2008	435	68.4	60	364.2	67.2	51	12.5%	70.80 [45.51, 96.09]	<del>-</del>
perwaiz 2009	442	32	53	370	38.5	55	13.8%	72.00 [58.67, 85.33]	
Subtotal (95% CI)			177			168	51.7%	47.28 [17.58, 76.98]	
Heterogeneity: $tau^2 =$	797.47;	chi <sup>2</sup> =	24.87,	df = 3 (1	P < 0.0	001); 1	$t^2 = 88\%$		
Test for overall effect	Z = 3.12	2 (P =	0.002)						
Total (95% CI)			448			445	100.0%	31.65 [7.14, 56.17]	
Heterogeneity: tau <sup>2</sup> =	1088.61;	chi <sup>2</sup> =	= 96.20	, df = 7	(P < 0)	.00001	); $I^2 = 93\%$	3	
Test for overall effect	Z = 2.53	3 (P =	0.01)					-100 F	-50 0 50 1
Test for subgroup dif	ferences:	chi <sup>2</sup> =	5.86, 0	df = 1 (P)	0 = 0.0	2), $I^2 =$	82.9%	Favo	ours [Roux-en-Y] Favours [Billroth II]

FIGURE 11: Forest plot of operation time.

analysis. Odds ratio (OR), mean difference (MD), and 95% confidence interval (CI) data were generated, and differences were considered significant when P < 0.05. Regarding heterogeneity assessment, a fixed-effects model was used when P > 0.1 and  $I^2 < 50\%$ , and a random-effects model was used when P < 0.1 and  $I^2 > 50\%$ . Subgroup analyses were performed when P < 0.1 and  $I^2 > 50\%$  to determine the reason for the heterogeneity. We prepared funnel plots to assess publication bias.

# 3. Results

- 3.1. Literature Search Results. Ultimately, nine high-quality articles [5–8, 13–17], including four RCTs [5–8] and five CCTs [13–17], were included. The literature screening process is shown in Figure 1. The general parameters of the included studies are shown in Table 1, and the pathologies, surgery-related parameters, and quality scores of the included studies are shown in Table 2.
- 3.2. Primary Outcomes. Nine studies [5–8, 13–17] provided the incidence of DGE, three [6, 7, 16] provided the incidence of A-grade DGE, four [5–7, 16] provided the incidence of B-grade DGE, and four [5–7, 16] provided the incidence of C-grade DGE. The heterogeneity results were P = 0.0002,  $I^2 = 74\%$ ; P = 0.03,  $I^2 = 70\%$ ; P = 0.32,  $I^2 = 14\%$ ; and P = 0.10,  $I^2 = 52\%$ , respectively, as shown in Figures 2–5. The fixed-effects model was selected for B-grade DGE, while the random-effects model was used for the other outcomes. The overall effect sizes of the above outcomes were as follows:

OR = 1.01, 95% CI: 0.50–2.03; OR = 0.49, 95% CI: 0.17–1.45; OR = 0.63, 95% CI: 0.29–1.38; and OR = 2.13, 95% CI: 0.38–11.99, respectively. These results suggest that the outcomes were not significantly different (P > 0.05) between the Roux-en-Y and Billroth II groups.

3.3. Secondary Outcomes. Nine studies [5-8, 13-17] provided the incidence of POPF, five [6-8, 14, 16] provided the incidence of abscess, five [6-8, 14, 16] provided the incidence of bile leaks, seven [7, 8, 13-17] provided the incidence of infection, five [6, 7, 13–15] provided the incidence of postoperative bleeding, eight [5-8, 13-15, 17] provided the operation time, and eight [5-8, 13-15, 17] provided the length of postoperative hospital stay. The heterogeneity test results were P = 0.92,  $I^2 = 0\%$ ; P = 0.65,  $I^2 = 0\%$ ; P = 0.38,  $I^2 = 4\%$ ; P = 0.45,  $I^2 = 0\%$ ; P = 0.59,  $I^2 = 0\%$ ; P < 0.00001,  $I^2 = 93\%$ ; and P < 0.0001,  $I^2 = 77\%$ , respectively, as shown in Figures 6-12. The random-effects model was used to analyze the length of operation and postoperative hospital stay data; for analysis of the other outcomes, the fixed-effects model was used. The overall effect sizes of the above outcomes were as follows: OR = 0.84, 95% CI: 0.62-1.15; OR = 1.22, 95% CI: 0.76-1.96; OR = 0.87, 95% CI: 0.48-1.58; OR = 1.10, 95% CI: 0.76-1.61; OR = 1.37, 95% CI: 0.64-2.95; MD = 31.65, 95% CI: 7.14-56.17; and MD = -0.72, 95% CI: -2.69-1.25, respectively. The results suggested no significant differences in outcomes (all P > 0.05) between the Roux-en-Y and Billroth groups, except for operation time (P < 0.05).

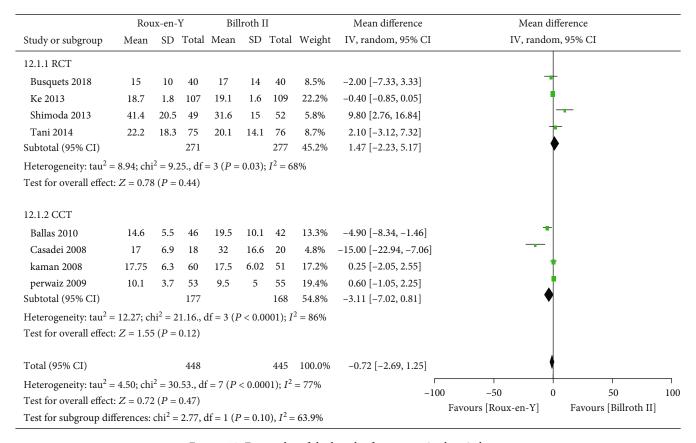


Figure 12: Forest plot of the length of postoperative hospital stay.

3.4. Sensitivity and Subgroup Analyses. The heterogeneity data for DGE were P = 0.0002,  $I^2 = 74\%$ ; the results did not change after the subgroup analysis, indicating high reliability thereof. The heterogeneity values of A-grade DGE and Cgrade DGE were P = 0.03,  $I^2 = 70\%$ , and P = 0.10,  $I^2 = 52\%$ , respectively. We eliminated each study one by one, and the results showed good stability. The heterogeneity data for operation time and length of postoperative hospital stay were P < 0.00001,  $I^2 = 93\%$ , and P < 0.0001,  $I^2 = 77\%$ , respectively. We conducted a subgroup analysis based on the RCT and CCT groups. Operation time in the RCT group was not significantly different between the two reconstructions, while in the CCT group Roux-en-Y reconstruction had a longer operation time (MD = 47.28, 95% CI: 17.58-76.98, P =0.002). Subgroup analysis did not reveal a significant difference in length of postoperative hospital stay between the Roux-en-Y group and Billroth II groups (P = 0.44 and P =0.12 for RCT and CCT, respectively). Therefore, operation time was associated with study type. A high-quality RCT or CCT is still needed to verify the operation time difference between the two reconstructions.

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3.5. Bias Analysis. We drew funnel plots based on the POPF data. The results showed that the 95% CI data were similar among the studies, which were distributed in a roughly symmetrical manner between the two sides of the midline, suggesting that the results were unaffected by publication bias and were thus highly reliable (Figure 13).

#### 4. Discussion

PD is a classic operation for benign and malignant lesions around the head of the pancreas. The incidence and complexity of postoperative complications are high due to the loss of organs and tissues [2, 21, 22]. DGE is a common complication after PD [23, 24]. Among the factors affecting DGE, the method used to reconstruct the digestive tract has been controversial. The use of Billroth II or Roux-en-Y reconstruction following PD to prevent DGE is also controversial [5–8, 13–17]. Meta-analyses that investigated the two reconstruction methods could not reach an agreement regarding which was superior [9, 10].

Consequently, we conducted this meta-analysis to analyze the characteristics of the two reconstruction methods and provide evidence-based guidance for clinical work. The results showed that traditional Billroth II reconstruction shortened the operation time compared to Roux-en-Y reconstruction, but no significant differences in any other complications were observed. Our results also indicated no significant differences between Roux-en-Y reconstruction and Billroth II reconstruction in DGE, A-grade DGE, B-grade DGE, or C-grade DGE (OR = 1.01, 95% CI: 0.50–2.03, P = 0.98; OR = 0.49, 95% CI: 0.17–1.45, P = 0.20; OR = 0.63, 95% CI: 0.29–1.38, P = 0.25; and OR = 2.13, 95% CI: 0.38–11.99, P = 0.39). Furthermore, no differences were detected in POPF, abscess, bile leak, infection, postoperative bleeding, or postoperative hospital stay (P > 0.05).

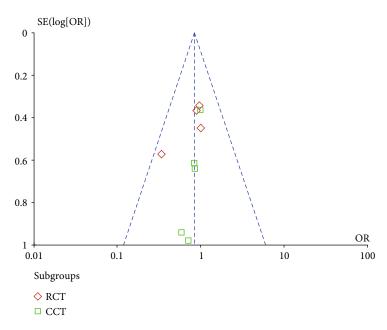


FIGURE 13: Funnel plot of the literature data on postoperative pancreatic fistula (POPF).

In 2015, Yang et al. [9] included three high-quality RCTs in their meta-analysis and reported that Billroth II reconstruction lowered the incidence of B- and C-grade DGE, although the small number of included studies was a limitation. Li et al. [10] conducted a case-control study that included 43 patients undergoing Roux-en-Y and 43 patients undergoing traditional Billroth II reconstruction after PD. The operation time of the Roux-en-Y group was longer than that of the traditional reconstruction group. A subsequent systematic review evaluated a series of postoperative complications. Klaiber et al. [25] systematically evaluated the postoperative complications of the two methods, and the results were largely consistent with those of the present study. However, it remains to be determined whether carrying out pancreatogastrostomy after PD had an impact on the overall results [26].

Based on previous studies, our study systematically screened the literature for studies of PD and excluded those in which PD was followed by pancreatogastrostomy. Articles with high heterogeneity were also excluded, so our analyses were characterized by relatively high homogeneity. We conducted this new systematic evaluation to overcome the deficiencies of previous research, and the results are reliable. However, this study also had some shortcomings. Although homogeneity was high, the number of included studies was small; hence, a larger study is needed.

We performed subgroup and sensitivity analyses of the outcomes with high heterogeneity; the results remained unchanged, except for operation time, further indicating high reliability. We also performed a subgroup analysis of operation time based on the RCTs and CCTs, and the results indicated that study type was the primary influencing factor. Therefore, an RCT or CCT with a large sample size is needed to further compare the operation times of Roux-en-Y and Billroth II digestive tract reconstruction.

In conclusion, we demonstrated that Roux-en-Y reconstruction took longer than Billroth II reconstruction after PD. However, complications were not different between the two reconstruction types. Therefore, it is suggested that consideration of the difference in operation time and patients' condition is needed to ensure that a suitable personalized surgical plan is implemented.

#### **Data Availability**

The datasets generated or analyzed during the current study are available from the corresponding author on reasonable request.

#### **Conflicts of Interest**

The authors declare that they have no conflicts of interest.

## **Authors' Contributions**

Fulin Ma and Yong Fan contributed equally to this study.

#### Acknowledgments

This work was supported by the Key Research and Development Project of Gansu Province (no. 20YF3FA033) and the Cuiyin Science and Technology Innovation Program of Lanzhou University Second Hospital (no. CY2017-BJ04).

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