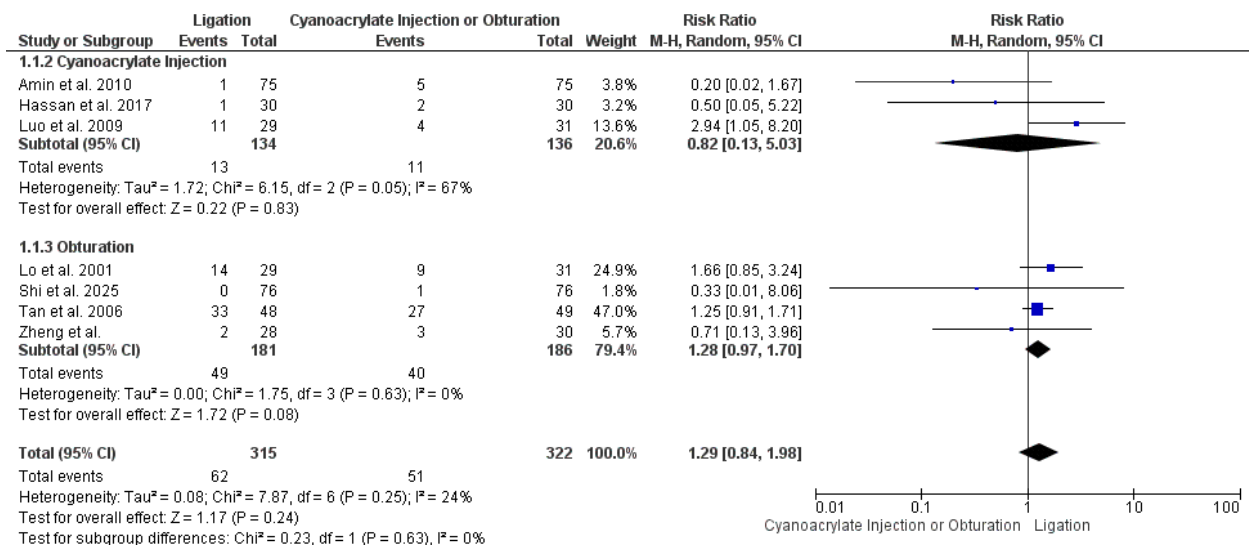
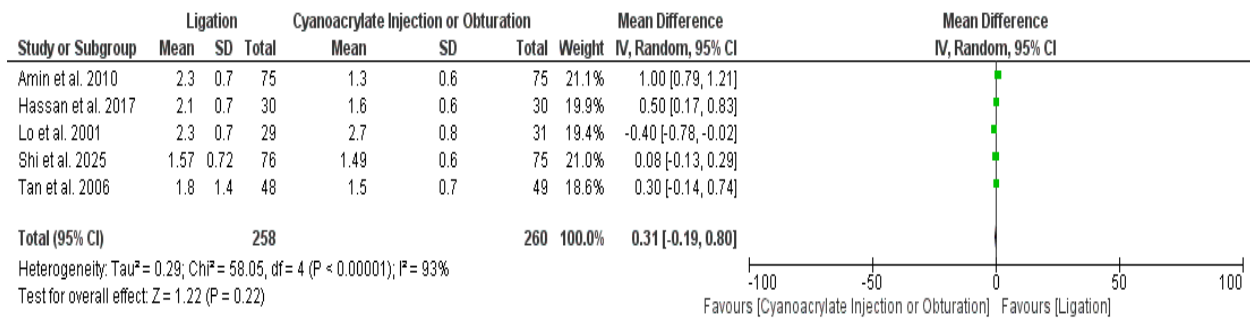


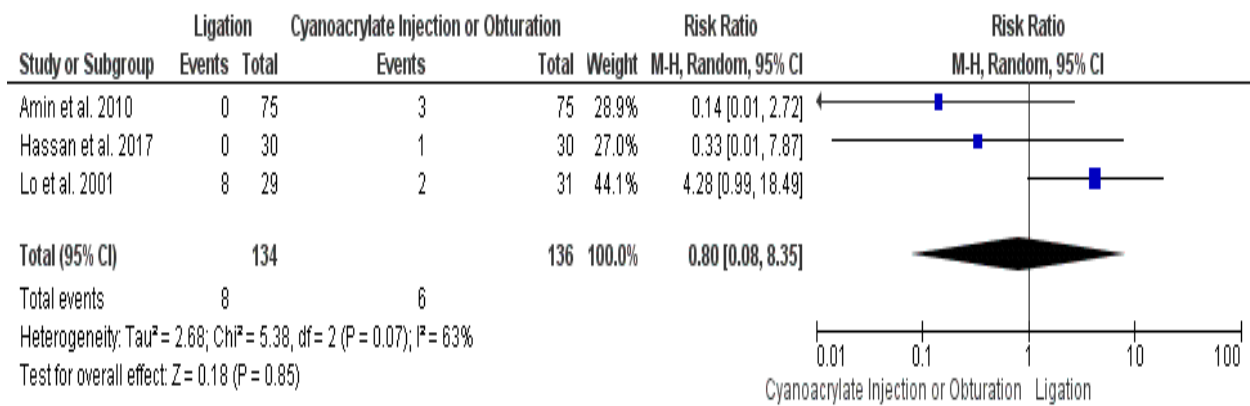
Supplementary Figure 1 Forest plot comparing all-cause mortality between EVL and cyanoacrylate therapy. Pooled risk ratio (RR) with 95% CI calculated using a random-effects model. No statistically significant difference was observed between groups (RR = 1.29; 95% CI [0.84–1.98]; I² = 21%). aMortality assessed at longest follow-up time.



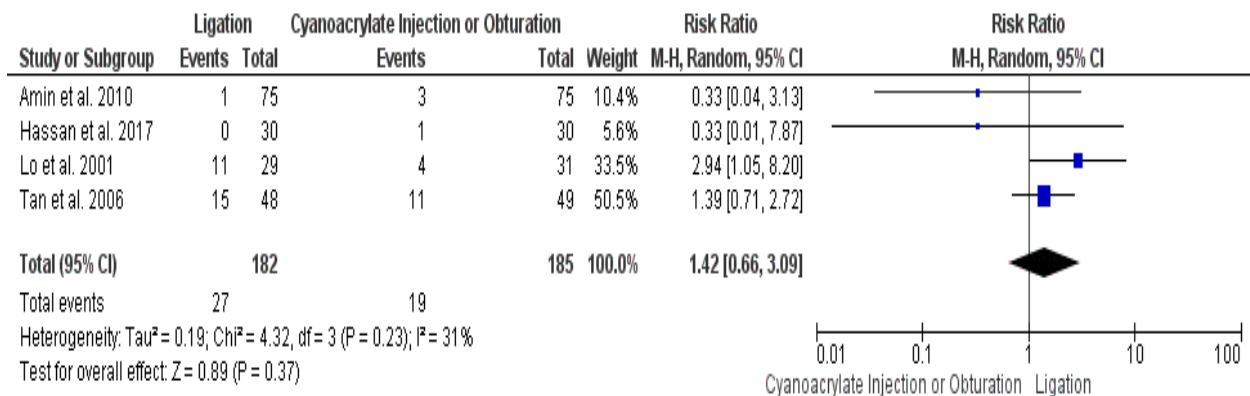
Supplementary Figure 2 Subgroup analysis of all-cause mortality by cyanoacrylate delivery method. Injection subgroup: RR = 0.82 (95% CI [0.13–5.03]); high heterogeneity (I² ≈ 70%). Obturation subgroup: RR = 1.28 (95% CI [0.97–1.70]); I² = 0%. Test for subgroup differences was non-significant (p > 0.05). a Subgroup analysis pre-specified in protocol.



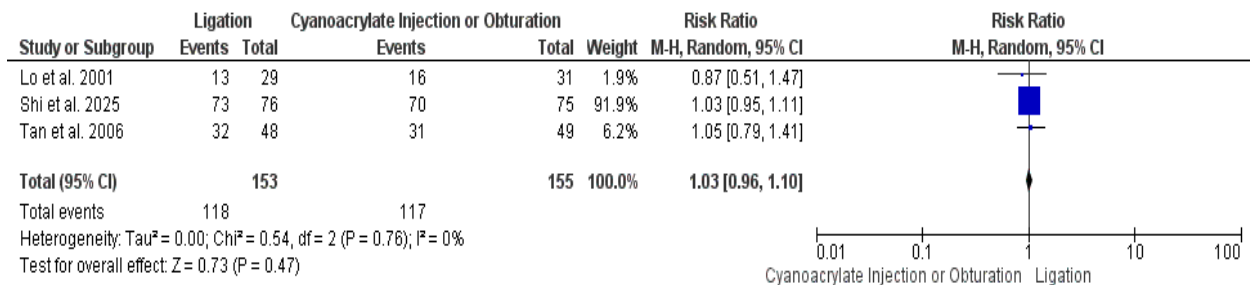
Supplementary Figure 3 Forest plot comparing the number of sessions required for variceal obliteration between EVL and cyanoacrylate therapy. No significant difference observed (MD = 0.31 sessions; 95% CI [-0.19–0.80]; I² = 76%). a MD = mean difference.



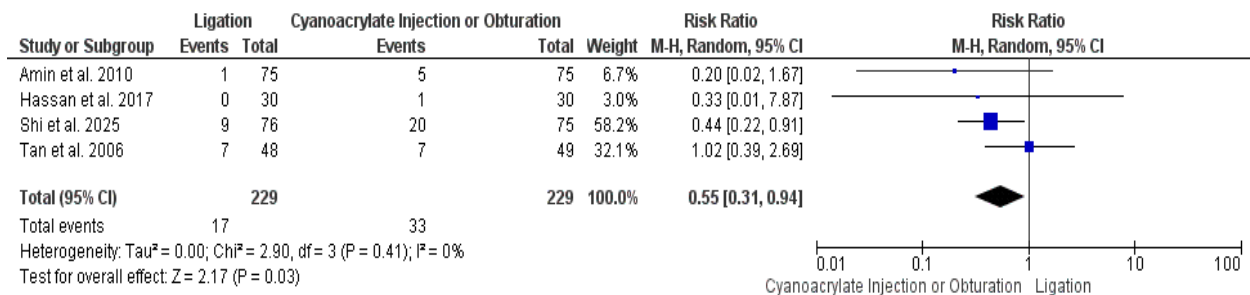
Supplementary Figure 4 Forest plot comparing fatal bleeding events between EVL and cyanoacrylate therapy. No significant difference observed between groups (RR = 0.80; 95% CI [0.08–8.35]; I² = 54%). a Fatal bleeding defined as death directly related to variceal hemorrhage.



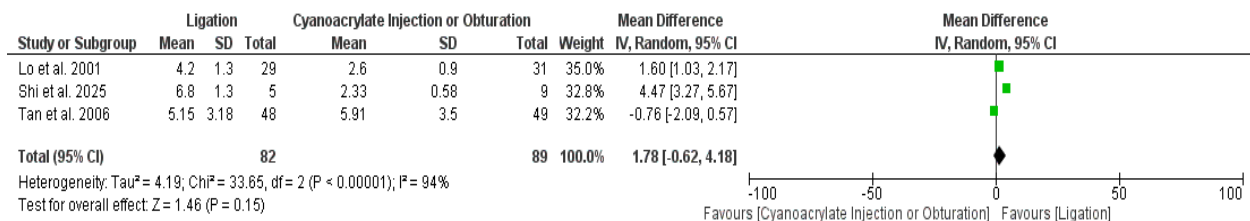
Supplementary Figure 5 Forest plot comparing treatment failure between EVL and cyanoacrylate therapy. No significant difference in treatment failure rates (RR = 1.42; 95% CI [0.66–3.09]; I² = 20%). a Treatment failure defined as failure to achieve initial hemostasis or early rebleeding.



Supplementary Figure 6 Forest plot comparing variceal obliteration rates between EVL and cyanoacrylate therapy. Pooled RR indicated no significant difference between interventions (RR = 1.03; 95% CI [0.96–1.10]; I² = 0%). a Obliteration confirmed endoscopically.



Supplementary Figure 7 Forest plot comparing post-procedural fever between EVL and cyanoacrylate therapy. EVL was associated with a significantly lower risk of post-procedural fever (RR = 0.55; 95% CI [0.31-0.94]; $I^2 = 0\%$). a Fever defined as $\geq 38^\circ\text{C}$ within 24 hours post-procedure.



Supplementary Figure 8 Forest plot comparing blood transfusion requirements between EVL and cyanoacrylate therapy. No significant difference observed (MD = 1.78 units; 95% CI not significant), with substantial heterogeneity ($I^2 > 75\%$). a Blood transfusion recorded during index hospitalization.

Supplementary Table 1 PICO and Search Strings

Population	Intervention	Control	Outcomes
Patients with gastric Varices	Endoscopic band ligation	Endoscopic Obturation/Cyanoacrylate Injection	Variceal Initial hemostasis/rebleeding
KeyWords			
PubMed	(((Cyanoacrylate Injection) OR (Cyanoacrylate)) AND (Band Ligation)) AND (gastric variceal bleeding)		78
Embase	('cyanoacrylate injection'/exp OR 'cyanoacrylate'/exp OR cyanoacrylate:ti,ab,kw OR 'cyanoacrylate injection':ti,ab,kw OR 'glue injection':ti,ab,kw) AND ('band ligation'/exp OR 'endoscopic band ligation'/exp OR 'endoscopic variceal ligation'/exp OR 'band ligation':ti,ab,kw OR 'endoscopic band ligation':ti,ab,kw OR 'endoscopic variceal ligation':ti,ab,kw OR evl:ti,ab,kw) AND ('gastric varices'/exp OR 'gastric variceal bleeding'/exp OR 'gastric variceal hemorrhage':ti,ab,kw OR 'bleeding gastric varices':ti,ab,kw OR 'gastric varices':ti,ab,kw)		43

Cochrane (Cyanoacrylate:ti,ab,kw OR "Cyanoacrylate 39
Injection":ti,ab,kw OR "Glue
Injection":ti,ab,kw) AND ("Band
Ligation":ti,ab,kw OR "Endoscopic Band
Ligation":ti,ab,kw OR "Endoscopic Variceal
Ligation":ti,ab,kw OR EVL:ti,ab,kw) AND
("Gastric Varices":ti,ab,kw OR "Gastric Variceal
Bleeding":ti,ab,kw OR "Gastric Variceal
Hemorrhage":ti,ab,kw OR "Bleeding Gastric
Varices":ti,ab,kw)

Clinical
Trials
Registry

5

Supplementary Table 2 Baseline characteristics of included studies

Stud y ID	Group	N	Age (me an ± SD)	Sex (M/ F)	HCC Prese nce	Albu min (g/dL) (mea n ± SD)	Biliru bin (mg/d L) (mean ± SD)	Hemogl obin (g/dL) (mean ± SD)	Platel ets (/mm ³) (mea n ± SD)	Creati nine (mg/d L) (mean ± SD)
Ami n (201 0)	EVL Cyanoacr ylate	7 5	52 ± 6	53/ 22	NR	3.38 ± 0.6	2.3 ± 0.8	8.5 ± 1.2	239 ± 71.2	0.98 ± 0.67
		7 5	51 ± 4	55/ 20	NR	3.0 ± 0.6	2.7 ± 1.0	8.4 ± 1.7	213.6 ± 81.2	1.1 ± 0.39
Hass an	EVL	3 0	49 ± 5	19/ 11	0/30	3.28 ± 0.4	2.1 ± 0.7	8.7 ± 1.3	230 ± 61.2	0.94 ± 0.52

(2017)	Cyanoacrylate	30	51 ± 3	21/9	0/30	3.12 ± 0.6	2.5 ± 0.9	8.3 ± 1.6	215 ± 75.3	1.0 ± 0.42
Lo (2001)	GVL	29	55 ± 13	22/7	10/29	3.2 ± 1.2	2.0 ± 1.0	NR	NR	NR
	GVO	31	58 ± 17	24/7	7/31	3.0 ± 1.1	2.3 ± 0.9	NR	NR	NR
Luo (2009)	EVL	29	51.2 ± 9	NR	NR	NR	NR	NR	NR	NR
	Cyanoacrylate	31	50.6 ± 3	NR	NR	NR	NR	NR	NR	NR
Shi (2025)	EBL	77	57.4 ± 9	58/19	16/77	NR	NR	NR	NR	NR
	EVO	77	55.3 ± 2	59/18	14/77	NR	NR	NR	NR	NR
Tan (2006)	GVL	48	61.7 ± 7	34/12.3	23/48	2.89 ± 0.54	1.93 ± 1.36	8.98 ± 2.29	90.56 ± 55.20	1.19 ± 0.68
	GVO	49	61.3 ± 5	35/14.6	23/49	2.90 ± 0.47	2.42 ± 3.37	9.18 ± 1.87	105.9 ± 67.87	1.37 ± 1.12

Zheng (etal.)	GVL	28	NR	NR	NR	NR	NR	NR	NR	NR
	GVO	30	NR	NR	NR	NR	NR	NR	NR	NR

Study ID)	Group	Prothrombin (mean \pm SD)	Ascites	Encephalopathy	GV Type (Sarin)	GV Size (Hashizume)	Concomitant		Hemodynamic	
							mitis	EV	Bleeding	Instability
Amidon (2010)	EVL	66.8 \pm 8.2	NR	0/75	GOV1	NR	NR	75/75	6	NR
	Cyanoacrylate	60.7 \pm 12.1	NR	0/75	GOV1	NR	NR	75/75	14	NR
Hassan (2017)	EVL	63.2 \pm 8.2	NR	0/30	NR	NR	NR	30/30	4	NR
	Cyanoacrylate	59.7 \pm 10.6	NR	0/30	NR	NR	NR	30/30	8	NR
					GOV1 =20,					
					GOV2 =8,	F1=2,				
			17 (59%)		IGV1=	F2=19,	28 (96%)			
	GVL	NR)	0/29	1	F3=8)	29/29	NR	4.2 \pm 1.3
					GOV1 =21,	F1=1,				
Lo (2001)			15 (48%)		GOV2 =6,	F2=21,	27 (87%)			
	GVO	NR)	0/31	=6,	F3=9)	31/31	NR	2.6 \pm 0.9

					IGV1=						
					4						
Luo	EVL	NR	NR	NR	NR	NR	NR	29/29	NR	NR	
(200	Cyanoacr										
9)	ylate	NR	NR	NR	NR	NR	NR	31/31	NR	NR	
					GOV1						
					=37,						
					GOV2						
					=24,	F1=8,					
					IGV1=	F2=47,					
	EBL	NR	NR	NR	16	F3=22	NR	77/77	30	NR	
					GOV1						
					=46,						
					GOV2						
Shi					=23,	F1=5,					
(202					IGV1=	F2=40,					
5)	EVO	NR	NR	NR	8	F3=32	NR	77/77	38	NR	
					GOV1						
					=26,						
					GOV2						
					=16,	F1=10,					
		2.63	± 26/4		IGV1=	F2=24,				5.15	±
	GVL	2.26	8	6/48	6	F3=14	NR	48/48	NR	3.18	
					GOV1						
					=27,	F1=10,					
Tan					GOV2	F2=22,				5.91	±
(200	GVO	2.44	± 25/4		=9,	F3=17	NR	49/49	NR	3.50	
6)	(Control)	1.55	9	5/49							

IGV1=

13

Zhe

ng GVL

(et (Interven

al.) tion) NR NR NR NR NR NR NR 28/28 NR NR

GVO

(Control) NR NR NR NR NR NR NR 30/30 NR NR

Values are expressed as mean ± SD or n (%), unless otherwise stated. NR = Not reported; EVL = Endoscopic Variceal Ligation; EBL = Endoscopic Band Ligation; GVL = Gastric Variceal Ligation; GVO = Gastric Variceal Obliteration; EVO = Endoscopic Variceal Obliteration; HCC = Hepatocellular carcinoma; GV = Gastric varices; EV = Esophageal varices; GOV = Gastroesophageal varices; IGV = Isolated gastric varices. GV type classified according to Sarin; GV size according to Hashizume.

Supplementary Table 3 Study characteristics of included trials

Study ID	Country	Study Design	Population (Sample Size, Age, Sex)	Intervention (Type, Dose, Duration)	Comparator (if any)	Main inclusion criteria	Primary Outcome	Secondary outcome	Conclusion
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A mi n et al. (20 10)	Cair o, Egyp t; cond ucte d at three endo scop y cente rs in uppe r Egyp t: Assi ut, Min ya and Al- azha r univ ersit y	A rando mize d Rand omize d contr ol clinic al trial Sex: 108 males and 42 femal es (72% males, 28% femal es)	150 patien ts. Ag e: range d from 20 to 60 years, mean ± SD of 52.0 ± 6.0 years. Sex: 108 males and 42 femal es (72% males, 28% femal es)	EVL of juncti onal varice s (Grou p I: 75 patien ts). Sessio ns contin ued till obliter ation of the varice s. Repea ted every 2 weeks until obliter ation	Cyanoacry late injection (Group II: 75 patients). N-butyl-2- cyanoacryl ate (Histoacry l Blue, Germany) injected in travariceal ly (0.5 mL cyanoacryl ate + 0.7 mL Lipiodol per aliquot). Patients with active bleeding from junctiona l varices	Patients with portal hyperten sion based on clinical, laborator y and imaging studies; Patients with clinical signs of hemate mesis, melena or hematoc hezia; Patients with bleeding from junctiona l varices	Initia l hem ostas is (defi ned as cessa tion of blee ding for more than 72 hour s).	Surviv al time and compl icatio ns or death.	EVL may be a good alterna tive to cyano acrylat e injecti on in treatm ent of bleedi ng junctio nal varices .
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	hospitals			achieved.		(active spurting or oozing); Patients who gave informed written consent; Aged 20 to 60 years.			
Ha	Servi	A	60	EVL	Cyanoacrylate	Patients	Initial	Survival	Cyanoacrylate
ssa	ces	rando	patients.	of	late	aged	1	and	glue
n	Hos	mize	Ag	gastric	injection	between	hem	and	injecti
et	pital,	d	e:	varice	(Group II:	20 and 70	ostas	compl	on is
al	Laho	contr	range	s	30	years;	is	icatio	on is
	re	olled	d	(Grou	patients).	endosco	(defi	ns or	superi
	(Imp	trial	from	p I: 30	Endoscopy-	py-	ned	death.	or to
	lied		25 to	patien	c sessions	proven	as		EVL
	Paki		72	ts).	continued	acute	cessa		for
	stan)		years,	Endos	till	GVH	tion		achiev
			with a	copic	obliteratio	(includin	of		ing
			mean	sessio	n of the	g clinical	blee		hemos
			\pm SD	ns	varices,	signs of	ding		tasis
			of 50.0	contin	repeated	bleeding,	for		and
			\pm 4.0	ued		endosco	more		preve

			years.	till	every	2	pic	than	ning
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			femal	repeat			a, or		reblee
			es	ed			large GV		ding
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				2			color		has no
				weeks			signs		advant
				.			and no		age
							other		over
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Lo	Kao	Prosp	60	Endos	Band	Patients	Initia	Reblee	Endos
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	ral	Trial	31,	31	-active	variceal	ned	ulcer	acrylat
	Hos		GVL:	patien	ligating	bleeding;	as	bleedi	e

pital, Taip ei, Taiw an, Rep ublic of Chin a	29). A ge: GVO 58 ± 17, GVL 55 ± 13. Se x: GVO 24/7 M/W, GVL 22/7 M/W. (rarely 4 mL) used per sessio n. Perfor med regula rly at 3- to 4- week interv	ts). N- butyl- 2- cyano acryla te (Histo acryl) 0.5 mL mixed with 1.5 mL Lipiod ol 2 mL n. n. Perfor med regula rly at 3- to 4- week interv	device used. 1 to 4 rubber bands applied during 1 session. Performed regularly at 3- to 4- week intervals until obliteratio n.	between 20 and 70 years of age. GV bleeding defined by active spurting /oozing or stigmata of recent hemorrh age.	cessa tion of blee ding for more than 72 hour s). requir ement s, and surviv al.	ng, bacter emia, bacter ial perito nitis), blood transf usion requir ement s, and surviv al.	prove d more effecti ve and safer than band ligatio n in the manag ement of bleedi ng gastric varices .
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RCT = Randomized controlled trial; ITT = Intention-to-treat; GVH = Gastric variceal hemorrhage; UGI = Upper gastrointestinal; EIS = Endoscopic Injection Sclerotherapy. Primary outcomes and inclusion criteria are presented as reported in the original publications.

Supplementary Table 4 Prevalence of Cirrhosis, Non-Cirrhotic Portal Hypertension, and Ascites in Included Studies

Study	Cirrhosis (%)	Non-Cirrhotic Hypertension (%)	Portal Ascites Present (%)
Lo <i>et al.</i> (2001)	100% (60/60)	0%	53.3% (32/60)
Tan <i>et al.</i> (2006)	100% (97/97)	0%	52.6% (51/97)
Zheng <i>et al.</i> (2006)	100% (58/58)	NR	NR
Amin <i>et al.</i> (2010)	100% (150/150)	0%	NR
Luo <i>et al.</i> (2011)	NR	NR	NR
Hassan <i>et al.</i> (2018)	100% (60/60)	0%	NR
Shi <i>et al.</i> (2025)	NR	NR	NR

Note: Data are presented as percentage (number/total) where available. **NR:** Not reported. **GOV:** Gastroesophageal varices; **IGV:** Isolated gastric varices. For **Zheng *et al.*** and **Hassan *et al.***, the study methods explicitly stated that all included patients had liver cirrhosis, though individual patient data were not tabulated. **Amin *et al.*** reported Child-Pugh scores implying the presence of ascites but did not provide the exact number of patients with ascites.