

World Journal of *Gastrointestinal Endoscopy*

World J Gastrointest Endosc 2023 March 16; 15(3): 84-194



REVIEW

- 84 Gastroesophageal reflux disease in children: What's new right now?

Sintusek P, Mutalib M, Thapar N

MINIREVIEWS

- 103 Endoscopic techniques for gastric neuroendocrine tumors: An update
Massironi S, Gallo C, Laffusa A, Ciuffini C, Conti CB, Barbaro F, Boskoski I, Dinelli ME, Invernizzi P
- 114 Endoscopic advances in the management of gastric cancer and premalignant gastric conditions
Park E, Nishimura M, Simoes P
- 122 Endoscopic ultrasound guided biliary drainage in surgically altered anatomy: A comprehensive review of various approaches
Sundaram S, Kale A
- 133 Quality of bowel preparation in patients with inflammatory bowel disease undergoing colonoscopy: What factors to consider?
Gravina AG, Pellegrino R, Romeo M, Palladino G, Cipullo M, Iadanza G, Olivieri S, Zagaria G, De Gennaro N, Santonastaso A, Romano M, Federico A

ORIGINAL ARTICLE

Case Control Study

- 146 Orientation in upper gastrointestinal endoscopy – the only way is up
Sivananthan A, Kerry G, Darzi A, Patel K, Patel N

Retrospective Study

- 153 Aluminum phosphate gel reduces early rebleeding in cirrhotic patients with gastric variceal bleeding treated with histoacryl injection therapy
Zeng HT, Zhang ZL, Lin XM, Peng MS, Wang LS, Xu ZL

Prospective Study

- 163 Medium-term surgical outcomes and health-related quality of life after laparoscopic *vs* open colorectal cancer resection: SF-36 health survey questionnaire
Hung CM, Hung KC, Shi HY, Su SB, Lee HM, Hsieh MC, Tseng CH, Lin SE, Chen CC, Tseng CM, Tsai YN, Chen CZ, Tsai JF, Chiu CC

META-ANALYSIS

- 177 Endoscopic biliary treatment of unresectable cholangiocarcinoma: A meta-analysis of survival outcomes and systematic review
Rebhun J, Shin CM, Siddiqui UD, Villa E

CASE REPORT

- 191** Colonic ductal adenocarcinoma case report: New entity or rare ectopic degeneration?

Conti CB, Mulinacci G, Tamini N, Jaconi M, Zucchini N

ABOUT COVER

Editorial Board Member of *World Journal of Gastrointestinal Endoscopy*, Pavel Skok, MD, PhD, Full Professor, Senior Adviser, Department of Gastroenterology, University Clinical Centre Maribor, Maribor 2000, Slovenia.
pavel.skok@ukc-mb.si

AIMS AND SCOPE

The primary aim of *World Journal of Gastrointestinal Endoscopy* (WJGE, *World J Gastrointest Endosc*) is to provide scholars and readers from various fields of gastrointestinal endoscopy with a platform to publish high-quality basic and clinical research articles and communicate their research findings online.

WJGE mainly publishes articles reporting research results and findings obtained in the field of gastrointestinal endoscopy and covering a wide range of topics including capsule endoscopy, colonoscopy, double-balloon enteroscopy, duodenoscopy, endoscopic retrograde cholangiopancreatography, endosonography, esophagoscopy, gastrointestinal endoscopy, gastroscopy, laparoscopy, natural orifice endoscopic surgery, proctoscopy, and sigmoidoscopy.

INDEXING/ABSTRACTING

The WJGE is now abstracted and indexed in Emerging Sources Citation Index (Web of Science), PubMed, PubMed Central, Reference Citation Analysis, China National Knowledge Infrastructure, China Science and Technology Journal Database, and Superstar Journals Database. The 2022 edition of Journal Citation Reports® cites the 2021 Journal Citation Indicator (JCI) for WJGE as 0.33.

RESPONSIBLE EDITORS FOR THIS ISSUE

Production Editor: Yi-Xuan Cai; Production Department Director: Xu Guo; Editorial Office Director: Jia-Ping Yan.

NAME OF JOURNAL

World Journal of Gastrointestinal Endoscopy

ISSN

ISSN 1948-5190 (online)

LAUNCH DATE

October 15, 2009

FREQUENCY

Monthly

EDITORS-IN-CHIEF

Anastasios Koulaouzidis, Bing Hu, Sang Chul Lee, Joo Young Cho

EDITORIAL BOARD MEMBERS

<https://www.wjgnet.com/1948-5190/editorialboard.htm>

PUBLICATION DATE

March 16, 2023

COPYRIGHT

© 2023 Baishideng Publishing Group Inc

INSTRUCTIONS TO AUTHORS

<https://www.wjgnet.com/bpg/gerinfo/204>

GUIDELINES FOR ETHICS DOCUMENTS

<https://www.wjgnet.com/bpg/GerInfo/287>

GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH

<https://www.wjgnet.com/bpg/gerinfo/240>

PUBLICATION ETHICS

<https://www.wjgnet.com/bpg/GerInfo/288>

PUBLICATION MISCONDUCT

<https://www.wjgnet.com/bpg/gerinfo/208>

ARTICLE PROCESSING CHARGE

<https://www.wjgnet.com/bpg/gerinfo/242>

STEPS FOR SUBMITTING MANUSCRIPTS

<https://www.wjgnet.com/bpg/GerInfo/239>

ONLINE SUBMISSION

<https://www.f6publishing.com>



Endoscopic biliary treatment of unresectable cholangiocarcinoma: A meta-analysis of survival outcomes and systematic review

Jeffrey Rebhun, Claire M Shin, Uzma D Siddiqui, Edward Villa

Specialty type: Gastroenterology and hepatology

Provenance and peer review: Unsolicited article; Externally peer reviewed.

Peer-review model: Single blind

Peer-review report's scientific quality classification

Grade A (Excellent): A
Grade B (Very good): 0
Grade C (Good): C
Grade D (Fair): 0
Grade E (Poor): 0

P-Reviewer: Koutsoumourakis A, Greece; Rahmati M, Iran

Received: November 3, 2022

Peer-review started: November 3, 2022

First decision: December 11, 2022

Revised: January 12, 2023

Accepted: March 1, 2023

Article in press: March 1, 2023

Published online: March 16, 2023



Jeffrey Rebhun, Department of Gastroenterology, Oregon Health and Sciences University, Portland, OR 97239, United States

Claire M Shin, Department of Medicine, University of Illinois at Chicago, Chicago, IL 60612, United States

Uzma D Siddiqui, Center for Endoscopic Research and Therapeutics, University of Chicago, University of Chicago Medicine, Chicago, IL 60637, United States

Edward Villa, Department of Gastroenterology and Hepatology, Northshore University Health System, Evanston, IL 60201, United States

Corresponding author: Jeffrey Rebhun, MD, Academic Fellow, Department of Gastroenterology, Oregon Health and Sciences University, 3181 SW Sam Jackson Park Road, Portland, OR 97239, United States. jeffrebbhun@gmail.com

Abstract

BACKGROUND

Endoscopic radiofrequency ablation (ERFA), percutaneous radiofrequency ablation (PRFA), and photodynamic therapy (PDT), when used in conjunction with conventional biliary stenting, have demonstrated a survival benefit in patients with unresectable cholangiocarcinoma.

AIM

To compare pooled survival outcomes, adverse event rates, and mean stent patency for those undergoing these procedures.

METHODS

A comprehensive literature review of published studies and abstracts from January 2011 to December 2020 was performed comparing survival outcomes in patients undergoing ERFA with stenting, biliary stenting alone, PRFA with stenting, and PDT with stenting for unresectable cholangiocarcinoma (CCA).

RESULTS

Data from four studies demonstrated a pooled mean survival favoring ERFA as compared to biliary stenting alone (12.0 ± 0.9 mo *vs* 6.8 ± 0.3 mo, $P < 0.001$) as well as statistically improved median survival time (13 mo *vs* 8 mo, $P < 0.001$). Both ERFA with stenting and PRFA with stenting groups demonstrated statistical superiority to biliary stenting alone ($P < 0.001$ and $P = 0.004$, respectively).

However, when comparing ERFA to PRFA, pooled data demonstrated overall higher mean survival in the ERFA with stenting cohort as compared to PRFA with stent cohort (12.0 ± 0.9 mo *vs* 8.1 ± 2.1 mo, $P < 0.0001$). Data from two studies demonstrated a pooled median survival favoring ERFA with stenting as compared to PDT with stenting (11.3 mo *vs* 8.5 mo, $P = 0.02$).

CONCLUSION

While further prospective, randomized studies are needed to assess efficacy of ERFA, our meta-analysis demonstrated that this technique offers endoscopists a reasonable palliative method by which to treat patients with unresectable CCA that results in longer survival as compared to biliary stenting alone, percutaneous radiofrequency ablation with biliary stenting, and PDT with biliary stenting as well as an acceptable adverse event profile based on available published data.

Key Words: Endoscopic radiofrequency ablation; Percutaneous radiofrequency ablation; Photodynamic therapy; Cholangiocarcinoma; Meta-analysis; Systematic review

©The Author(s) 2023. Published by Baishideng Publishing Group Inc. All rights reserved.

Core Tip: Endoscopic radiofrequency ablation offers endoscopists a reasonable palliative method by which to treat patients with unresectable cholangiocarcinoma that results in longer survival as compared to biliary stenting alone, percutaneous radiofrequency ablation, with biliary stenting, and photodynamic therapy with biliary stenting.

Citation: Rebhun J, Shin CM, Siddiqui UD, Villa E. Endoscopic biliary treatment of unresectable cholangiocarcinoma: A meta-analysis of survival outcomes and systematic review. *World J Gastrointest Endosc* 2023; 15(3): 177-190

URL: <https://www.wjgnet.com/1948-5190/full/v15/i3/177.htm>

DOI: <https://dx.doi.org/10.4253/wjge.v15.i3.177>

INTRODUCTION

Cholangiocarcinoma (CCA) is a primary cancer of the bile ducts accounting for 15% of primary hepatic malignancies and nearly 3% of malignant gastrointestinal tumors. 90% of CCA are extrahepatic (perihilar or main bile duct), while the remaining 10% are intrahepatic[1-3]. Due to location and delayed onset of symptoms, CCA has a poor prognosis with 5-year survival rates of 2%-25% and median survival of 3-6 mo for unresectable cancers[1,4]. 20%-30% of cholangiocarcinoma cases are surgically resectable, leaving the majority of CCA patients with only palliative options, namely, systemic chemotherapy and relief of biliary obstruction through surgical, percutaneous, and endoscopic approaches. The complex molecular landscape of cholangiocarcinoma, however, has limited the effectiveness of systemic chemotherapy in the treatment of unresectable cancer[5,6]. As a result of poor chemotherapeutic options, the mainstay of care for these patients with unresectable CCA revolves around endoscopic retrograde cholangiopancreatography (ERCP), interventional radiologic, or endoscopic ultrasound (EUS)-guided approaches for biliary decompression with biliary stenting and/or percutaneous drainage. While in the majority of cases these approaches are technically feasible and particularly effective at relieving biliary obstruction, the life-prolonging effects of these interventions remain poor, and adverse events, such as stent occlusion and cholangitis, limit their overall effectiveness [7,8].

Photodynamic therapy (PDT) is a well-studied, ablative technique resulting in cellular apoptosis or necrosis in cells that absorb a photosensitizer, an agent activated by a specific wavelength of light[9,10]. PDT protocols for CCA involve a two-stage treatment consisting of systemic administration of the photosensitizing agent (that is preferentially absorbed by pre-malignant and malignant tissue) followed 48 to 96 h later with transpapillary intra-biliary placement of a laser-emitting diode placed into the bile duct *via* cholangioscopy or ERCP. This diode, when activated, emits a wavelength of 630 nanometers (nm), and when directed towards cells that have absorbed the photosensitizer, results in cell death and necrosis of the target tissue. In a recent meta-analysis of ten studies assessing outcomes of PDT combined with biliary stenting compared to conventional biliary stenting alone, survival in the PDT group was 413 d, which was statistically superior to the 183 d for patients who underwent biliary stenting alone[10].

The limitations of this technology involve the two-stage approach and the resulting phototoxicity of the skin from the photosensitizer (lasting 4-6 wk in decreasing intensity), occurring in 0%-25% of patients undergoing PDT with meta-analytic data demonstrating a photosensitivity rate of 10.5%[9-15].

To minimize the risk of this adverse event, most protocols requires the patient to take significant measures to prevent any exposure to light following administration of the photosensitizer. Other reported adverse events reported include cholangitis and hepatic abscess.

Radiofrequency ablation (RFA) is a technology that delivers thermal energy *via* a catheter or probe to malignant tissue, resulting in locoregional coagulative necrosis and cellular death. RFA has been previously used successfully *via* percutaneous (PRFA) or intraoperative routes for the treatment of other solid organ tumors[16]. However, there is limited data available evaluating the role of endoscopic biliary RFA (ERFA) and PRFA as palliative measures in patients with unresectable cholangiocarcinoma. Our meta-analysis aims to evaluate survival outcomes of ERFA with biliary stenting compared with both the conventional stent-only approach and PRFA with stenting in the setting of unresectable CCA.

MATERIALS AND METHODS

A comprehensive literature search was conducted querying the PubMed, EMBASE, and Cochrane databases from January 2011 to December 2020. Keywords in our search included: “endoscopic radiofrequency ablation” and “cholangiocarcinoma”. In compiling studies assessing percutaneous radiofrequency ablation, the keywords in our search included: “percutaneous radiofrequency ablation” and “cholangiocarcinoma”. In compiling studies assessing photodynamic therapy, the keywords in our search included: “cholangiocarcinoma” and “photodynamic therapy.” The connector word “AND” was used to capture articles that were pertinent to our study. Reference articles were analyzed multiple authors for use in our initial inclusion. Our study was limited to articles published after the 2011 pilot study documenting the initial use of endoscopic radiofrequency ablation in human subjects[17]. Articles eligible for inclusion were limited to published retrospective (case-control studies) or prospective studies (randomized controlled trials) in the English language, conducted on human subjects. Additionally, studies included must have assessed both populations of interest with the intervention provided under similar medical conditions. Exclusion criteria included: Systematic reviews and/or meta-analyses; opinion papers; editorials; studies in which a contingency of data could not be extrapolated to generate the targeted outcome of survival duration; studies in which the patients underwent previous surgical intervention; studies in which other malignancies resulting in biliary obstruction (namely, pancreatic adenocarcinoma or ampullary carcinoma) were included, particularly if a contingency of data could not be extrapolated to generate the targeted conclusions or outcomes in cholangiocarcinoma subgroups. PRISMA flow charts (Figures 1A and B) were compiled to illustrate the results of our literature search with an additional detailed search strategy included as [Supplementary Table 1 and 2](#).

Three authors (Rebhun J, Shin CM, and Villa E) independently reviewed each article yielded from the above search strategy. Full text of the articles was then assessed to determine if inclusion criteria were met. Any missing or unclear data resulted in an attempt to contact the original author with relevant questions. Data pulled from each article included the following: Author and year of the article; Origin of the study; Type of study conducted; Subgroup total population; Patient age and gender distribution; Mean survival in months; Median survival in months; Mean stent patency in months; Adverse Events; Chemotherapy status.

Outcome assessed

Data was extracted from articles meeting inclusion criteria and combined to perform a meta-analysis. The primary objective was to compare mortality outcomes in patients undergoing endoscopic RFA with biliary stenting (henceforth to be referred to as the “ERFA” subgroup) to those undergoing endoscopic stenting alone as well as to those undergoing percutaneous RFA with biliary stenting (henceforth to be referred to as the “PRFA” subgroup). Secondary outcomes included duration of stent patency and rates of adverse events between the treatment groups.

Quality assessment of included studies

To better assess the quality of individual studies, we used the Newcastle-Ottawa scale (NOS) for retrospective case-control studies and the Cochrane tool for risk of bias for randomized controlled trials. The NOS uses 3 domains: Selection, comparability, and ascertainment of outcome to award a maximum of 9 total points. A score > 7 indicates a study of good quality. The NOS has been shown to be a marker of individual study quality when using non-randomized studies in meta-analyses[18,19]. NOS scores are reported in the supplementary portion of the article. In order to best evaluate the quality of evidence for each outcome amenable to meta-analysis, we used the Grading of Recommendations, Assessment, Development, and Evaluation system to interpret the clinical implications of our findings.

Statistical analysis

Continuous variables were reported as mean \pm standard deviation. Categorical variables were calculated as frequencies or percentages. Pooled survival data was used to generate Kaplan-Meier survival curves with log-rank test performed to assess for statistically significant differences in survival.

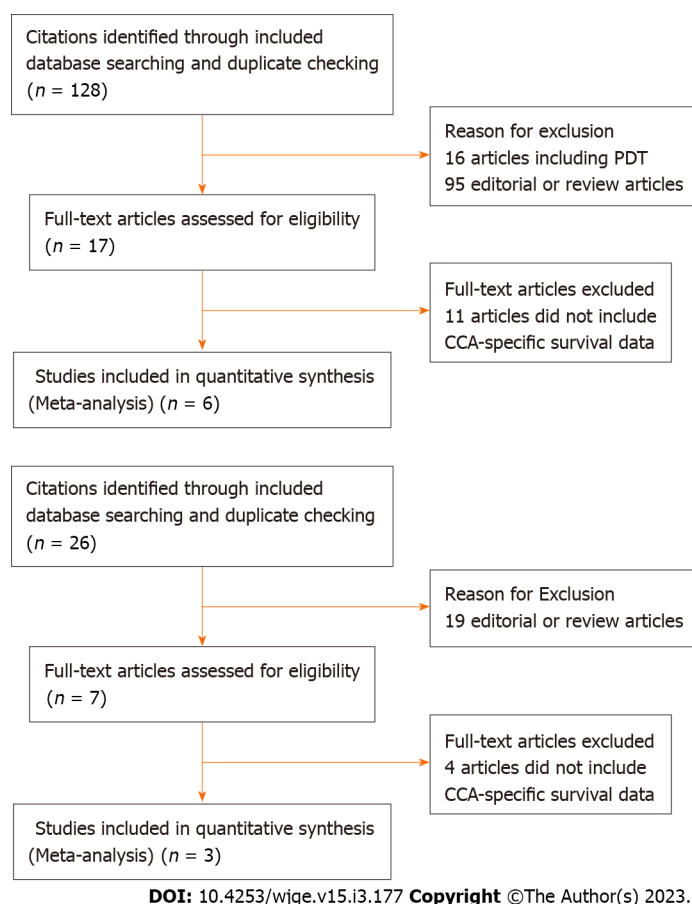


Figure 1 PRISMA flow charts. A: Flowsheet diagram demonstrating inclusion of studies for meta-analysis of endoscopic radiofrequency ablation (ERFA) with stenting versus biliary stenting alone; B: Flowsheet diagram demonstrating inclusion of studies for meta-analysis of ERFA with stenting versus photodynamic therapy with stenting. CCA: Cholangiocarcinoma; PDT: Photodynamic therapy.

Median days of survival was either reported in each study or extrapolated with use of study-specific survival tables and/or curves. Between-study heterogeneity was reported with the I^2 statistic with values greater than 50 suggestive of substantial heterogeneity[20]. Categorical data underwent chi-square analysis to ascertain statistically significant differences. Mann-Whitney U-Test was performed to compare mean stent patency. If survival or stent patency was reported in number of days, conversion to number of months was made by dividing number of days by 30.42. Time in months was then rounded to the nearest tenth decimal place. P values were 2-sided and statistical significance was achieved with a P value of < 0.05 . Data was analyzed using IBM SPSS Statistics for Macintosh, Version 27.0 (IBM Corp. Armonk, NY, United States). The datasets generated and/or analyzed during this study are available from the corresponding author on reasonable request.

RESULTS

ERFA compared to biliary stenting alone

Our initial search returned 128 studies. After exclusion of studies that did not satisfy inclusion criteria and/or met no exclusion criteria, four studies[21-24] were included for quantitative and qualitative analyses. Summary of study characteristics (Table 1) as well as procedural and survival outcomes of each study (Table 2) are demonstrated in the corresponding tables.

Patients in the ERFA cohort had a pooled mean survival time of 12.0 ± 0.9 mo ($I^2 = 37.0$) while patients undergoing stenting alone had a mean survival time of 6.8 ± 0.3 mo ($I^2 = 78.4$). Difference in survival was calculated to be 4.9 ± 0.1 mo and the analysis was associated with minimal heterogeneity ($P < 0.001$, $I^2 = 0$) (Figure 2). Median survival of the ERFA cohort was calculated to be 13 mo while median survival of the stent only cohort totaled 8 mo with log-rank test performed to suggest a significant difference ($P < 0.001$, Figure 3).

Two of four studies reported data on stent patency[21,22] (Table 3). Stent patency was not found to be significantly different in the study by Hu *et al*[21] ($P = 0.7$); however, stent patency was significantly higher in the ERFA cohort in Yang *et al*[22] ($P = 0.02$)[19,20]. Both studies contributed similarly to the

Table 1 Summary of included studies

Ref.	Country	Study type	Total patients	Mean age	Female gender (%)	Chemotherapy
Sampath <i>et al</i> [23], 2016	United States	Case-Control	25	69.7	10 (40.0)	19 (76)
Hu <i>et al</i> [21], 2016	China	RCT	63	71.4	32 (50.8)	-
Wu <i>et al</i> [26], 2017	China	Case-Control	71	57.9	28 (39.2)	59 (83)
Cui <i>et al</i> [25], 2017	China	Case-Control	39	64.7	17 (43.5)	2 (5)
Yang <i>et al</i> [22], 2018	China	RCT	65	63.2	32 (49.2)	-
Bokemeyer <i>et al</i> [24], 2019	Germany	Case-Control	44	67	-	13 (30)

RCT: Randomized controlled trial.

Table 2 Procedural and survival outcomes of individual studies

Ref.	Total patients		Technical success		Major adverse events		Mean survival (mean ± SD)		P value
	Stent only	RFA-stent	Stent only	RFA-stent	Stent only	RFA-stent	Stent only	RFA-stent	
Endoscopic									
Sampath <i>et al</i> [23], 2016	15	10	-	100	8	9	4.7 ± 5.5	12 ± 5.9	0.001
Hu <i>et al</i> [21], 2016	31	32	-	-	22	26	5.7 ± 0.5	10.4 ± 1.2	0.001
Yang <i>et al</i> [22], 2018	33	32	100	100	3	2	8.3 ± 0.5	13.2 ± 0.6	< 0.001
Bokemeyer <i>et al</i> [24], 2019	22	20	100	100	10	4	7.4 ± 0.9	11.4 ± 1.9	0.046
Percutaneous									
Wu <i>et al</i> [26], 2017	36	35	-	100	5	0	6.5 ± 2.6	8.4 ± 2.3	0.80
Cui <i>et al</i> [25], 2017	14	25	-	-	-	-	4.5 ± 2.1	6.7 ± 5.3	0.30

^aP value as it relates to mean survival in each respective study. SD: Standard deviation; RFA: Radiofrequency ablation.**Table 3 Pooled stent patency analysis among included endoscopic radiofrequency ablation studies**

Ref.	Stent only patients (%)	Mean stent patency	ERFA-stent patients (%)	Mean stent patency	P value
Hu <i>et al</i> [21], 2016	31 (48.5)	3.9	32 (50)	5	0.7
Yang <i>et al</i> [22], 2018	33 (51.5)	3.4	32 (50)	6.8	0.02
Cumulative	64	3.6	64	5.9	< 0.001

ERFA: Endoscopic radiofrequency ablation.

pooled analysis with only slightly more patients in the stent only treatment group being represented by Yang *et al*[22]. Pooled results of the two studies were calculated and demonstrated a mean stent patency in the ERFA with stent group to be 5.9 mo compared to 3.6 mo in the stent only group ($P < 0.001$). All four studies reported adverse event data and were used in our analysis (Table 4). Biliary stent occlusion was the most frequent adverse event that arose in both treatment groups, however there was no significant difference between ERFA (81%) and stent alone (67.3%, $P = 0.148$). Cholecystitis data was only reported in the Hu *et al*[21] and Bokemeyer *et al*[24] studies; however pooled analysis showed a 12.5% risk for cholecystitis in the ERFA cohort compared with 0% risk in the stent only cohort ($P = 0.01$). The frequency of hemobilia/bleeding was similar among the two groups (1.5% for both, $P = 1.0$).

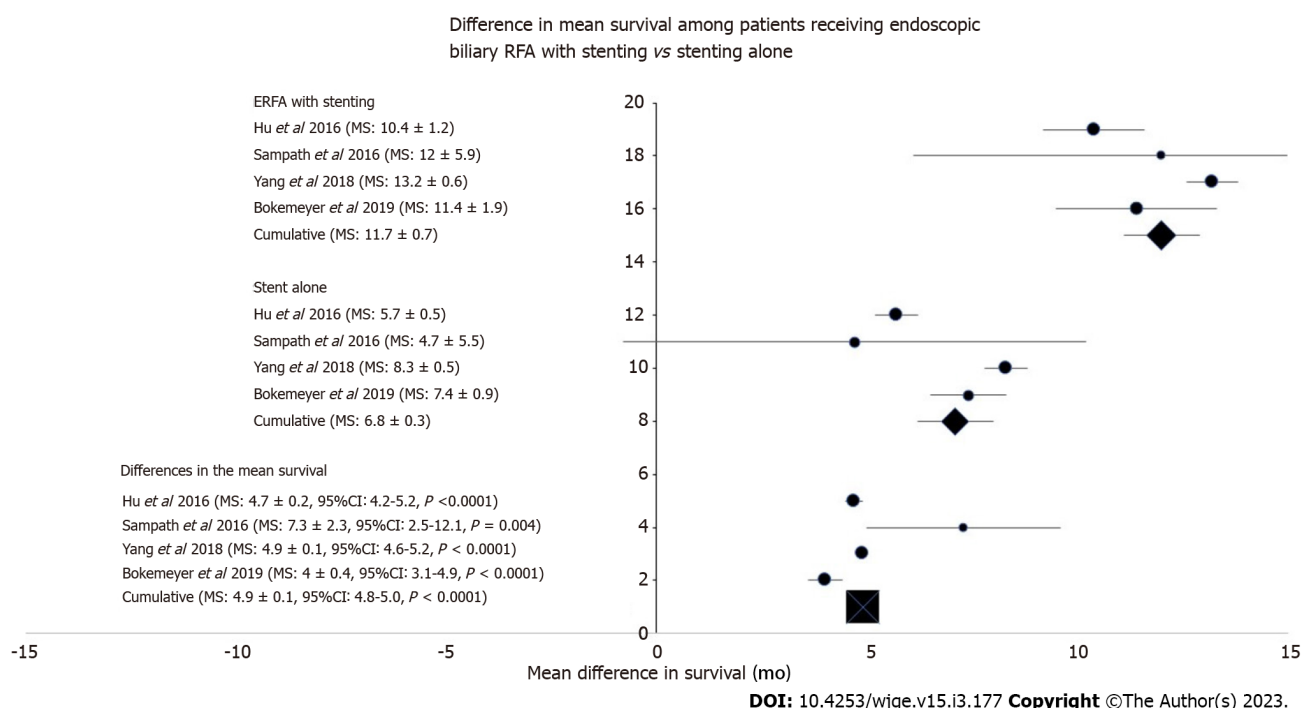
ERFA compared to percutaneous biliary RFA

Of the 128 articles in our initial literature search, six studies were included for meta-analytic comparisons of survival between ERFA and PRFA groups[21-26]. From these studies, there were 106 patients that underwent ERFA with concomitant stenting, and 60 patients who underwent PRFA with stenting for unresectable CCA. Comparison control groups included 101 patients who underwent

Table 4 Pooled adverse event data among included endoscopic radiofrequency ablation studies

Adverse event	ERFA-stent n (%)	Stent alone n (%)	P value
Biliary stent occlusion	34 (81.0)	31 (67.3)	0.148
Cholangitis	27 (25.5)	15 (19.0)	0.298
Cholecystitis	8 (12.5)	0 (0)	0.010
Pancreatitis	4 (4.2)	3 (4.7)	0.875
Hemobilia/Bleeding	1 (1.5)	1 (1.5)	1.000

ERFA: Endoscopic radiofrequency ablation.

**Figure 2 Forest plot of mean stent survival among treatment groups along with difference in survival. RFA: Radiofrequency ablation.**

biliary stenting in the ERFA studies and 50 patients who underwent biliary stenting in the PRFA studies.

The ERFA with stent cohort had a mean survival of 12.0 ± 0.9 mo ($Q = 4.8$, $I = 37\%$, Figure 4). The PRFA with stent cohort had a mean survival of 8.1 ± 2.1 mo ($Q = 0.09$, $I = 0\%$, Figure 4). In both ERFA and PRFA studies, mean survival was significantly increased compared to biliary stent alone control groups ($P < 0.001$ and $P = 0.004$, respectively). The difference in mean survival among both biliary RFA groups favored ERFA with stenting by 3.9 ± 0.2 mo (95%CI: 3.4-4.4, t -test = 16.6, $P < 0.0001$; Figure 4).

The ERFA group had a median survival (Figure 5) of 13 mo compared to the PRFA group median survival of 5.2 mo (log-rank test $Z = 5.3$, $P < 0.0001$). Only patients undergoing ERFA with stenting had a significant difference in median survival as compared to the biliary stent alone control group ($P < 0.001$).

Adverse event data went unreported in the Cu *et al* [25] study, thus comparison of PRFA adverse event was limited to those of procedures reported by Wu *et al* [26]. In comparing this study to those of the ERFA cohort, the risk of cholangitis was increased in the ERFA with stent cohort ($\chi^2 = 11.0$, $P = 0.001$).

ERFA compared to PDT

Of the 26 studies identified in our initial literature search, two studies provided data contingent for direct comparison of PDT and ERFA survival in patients with unresectable CCA (Table 5) [13,14]. From these studies, 49 patients underwent ERFA, and 56 underwent PDT (Table 5). All patients underwent concomitant biliary stenting whether *via* ERCP or *via* percutaneous transhepatic biliary drainage. Pooled median survival of the ERFA group was 11.3 mo, and median survival of the PDT group was 8.5 mo, a

Table 5 Demographics, procedural, and survival outcomes of endoscopic radiofrequency ablation vs photodynamic therapy

	Demographic data							
	Strand <i>et al</i> [13], 2014				Schmidt <i>et al</i> [15], 2016			
					Gao <i>et al</i> [14], 2018			
Number of patients	RFA	16		$P = 0.1$	RFA	14	NA	RFA
	PDT	32			PDT	20		PDT
Gender (male)	RFA	10		$P = 1.0$	RFA	8	$P = 0.1$	RFA
	PDT	19			PDT	6		PDT
Age (mean, yr)	RFA	64.3 ± 11.9		$P = 0.1$	RFA	73 ± 9	$P = 0.2$	RFA
	PDT	69.5 ± 13.6			PDT	70 ± 12		PDT
Number of treatments	RFA	28 (mean: 1.2)		$P = 0.02$	RFA	31	NA	RFA
	PDT	60 (mean: 2.1)			PDT	36		PDT
Median survival (month)	RFA	9.6		$P = 0.8$	RFA	NA	NA	RFA
	PDT	7.5			PDT	NA		PDT
Lead time to initial treatment (days)	RFA	NS		$P = 0.6$	RFA	300 ± 270	NA	RFA
	PDT	NS			PDT	120 ± 90		PDT
Total bilirubin concentration (μmol/dL)	RFA	NA		NA	RFA	3.3 ± 3.9	$P = 0.7$	RFA
	PDT	NA			PDT	4.1 ± 6.9		PDT
Tumor location	RFA	Intrahepatic	1	$P = 0.1$	RFA	Intrahepatic	1	$P = 0.5$
		Hilar	13			Hilar	11	
		Distal/Extrahepatic	2			Distal/Extrahepatic	1	
	PDT	Intrahepatic	0		PDT	Intrahepatic	3	PDT
		Hilar	32			Hilar	15	
		Distal/Extrahepatic	0			Distal/Extrahepatic	1	
N1 staging	RFA	7		$P = 0.8$	RFA	3	$P = 0.4$	RFA
	PDT	12			PDT	2		PDT
M1 staging	RFA	6		$P = 0.2$	RFA	2	$P = 0.8$	RFA
	PDT	6			PDT	6		PDT
Stents placed	Total	RFA	115	NA	Total	RFA	29	NA
		PDT	307			PDT	44	
	Plastic	RFA	69	NA	Plastic	RFA	26	NA
		PDT	264			PDT	38	
	Total metallic	RFA	46	NA	Total metallic	RFA	3	NA
		PDT	43			PDT	6	
	Fully covered	RFA	17	NA	Fully covered	RFA	NA	NA
		PDT	14			PDT	NA	
	Uncovered	RFA	29	NA	Uncovered	RFA	NA	NA
		PDT	29			PDT	NA	
Number of ERCPs	RFA	91		NA	RFA	NA	NA	RFA
	PDT	170			PDT	NA		PDT
Percutaneous transhepatic biliary drainage (PTBD)	RFA	2		$P = 0.2$	RFA	2	$P = 0.3$	RFA
	PDT	10			PDT	6		PDT

RFA: Radiofrequency ablation; PDT: Photodynamic therapy; NA: Not available; NS: Not significant; *P*: *P* value as it relates to each comparator category.

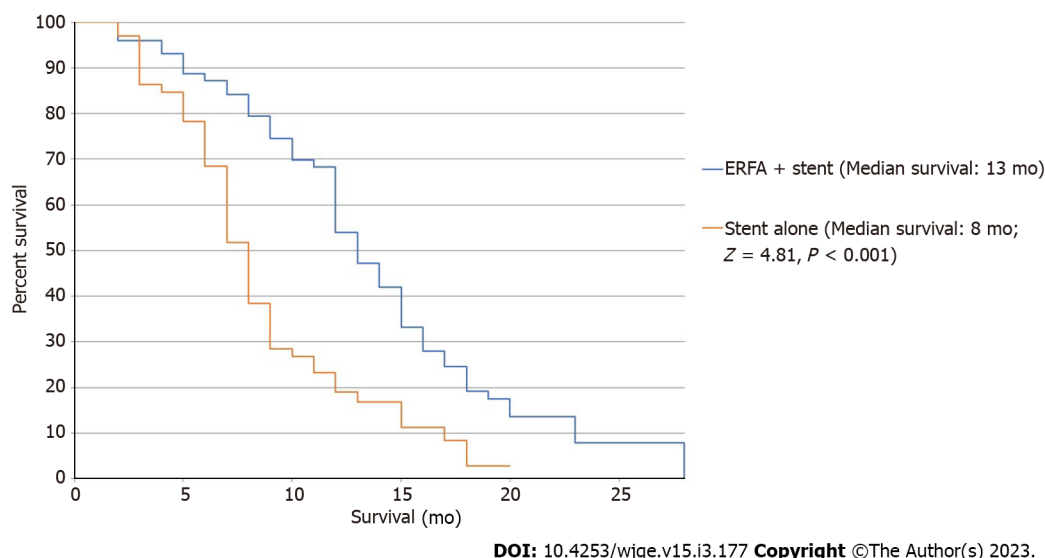


Figure 3 Kaplan-Meier survival curve of endoscopic radiofrequency ablation with stenting vs stenting alone. ERFA: Endoscopic radiofrequency ablation.

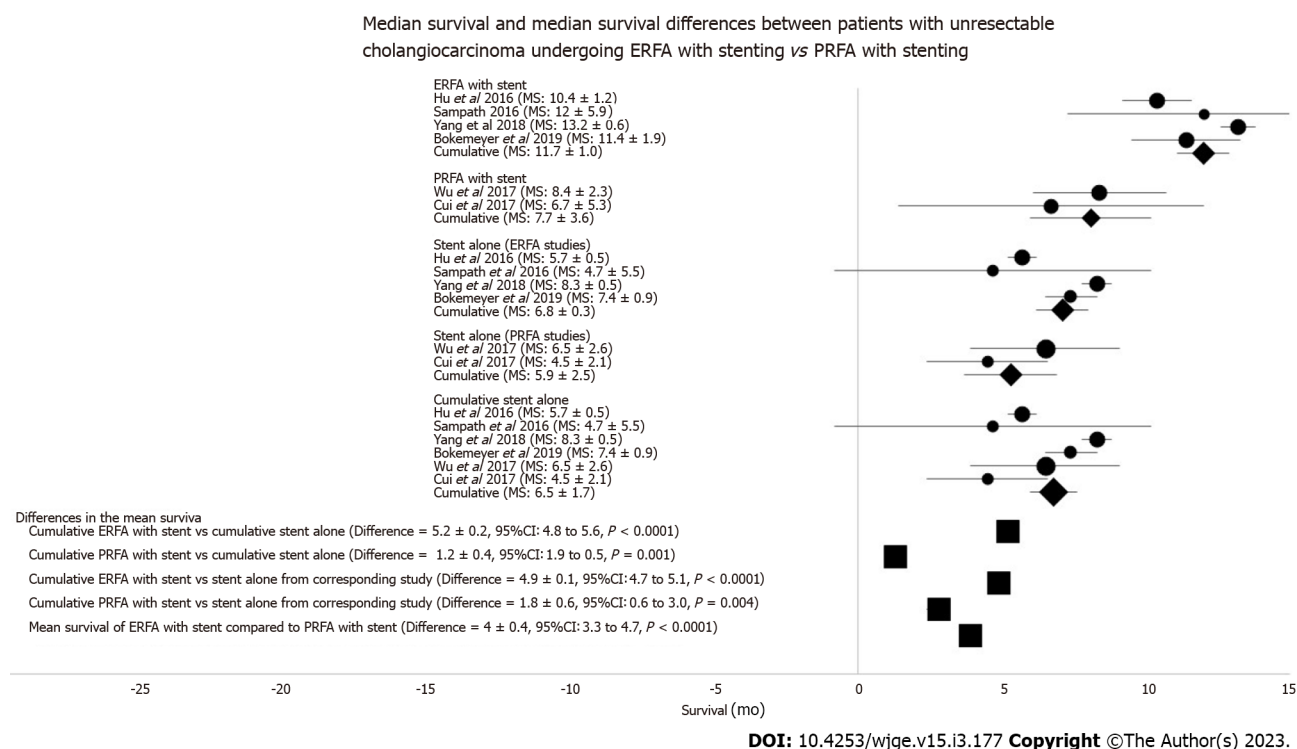


Figure 4 Forest plot of mean survival of endoscopic radiofrequency ablation with stenting; percutaneous radiofrequency ablation with stenting; comparisons to corresponding biliary stenting alone subgroups; and overall comparisons in mean survival. ERFA: Endoscopic radiofrequency ablation; PRFA: Percutaneous radiofrequency ablation.

difference that was statistically significant (Figure 6; *P* = 0.02).

Of the 26 studies identified, three studies provided data contingent for direct comparison of PDT and ERFA adverse events (Table 5)[13-15]. With regard to pooled adverse events among 62 patients who underwent ERFA and 75 patients who underwent PDT, there were statistically higher rates of stent occlusions (22.6% *vs* 6.7%, *P* = 0.008) and cholangitis (74% *vs* 41.3%, *P* = 0.001) in the ERFA group (Table 6); however, there were increased rates of stent migration (16% *vs* 4.8%, *P* = 0.04), moderate or

Table 6 Adverse events of endoscopic radiofrequency ablation vs photodynamic therapy

Adverse events	RFA	PDT	P value
Stent related complications	17	17	0.7
Stent occlusion	14	5	0.008
Stent migration	3	12	0.04
Cholangitis	46	31	0.001
Hepatic abscess	4	3	0.5
Bleeding	1	1	0.9
Moderate/Severe abdominal pain	3	17	0.003
Post-ERCP pancreatitis	3	2	0.5
Phototoxicity	0	2	NA

ERCP: Endoscopic retrograde cholangiopancreatography; NA: Not applicable; RFA: Radiofrequency ablation; PDT: Photodynamic therapy; P: P value as it relates to comparisons of each adverse event.

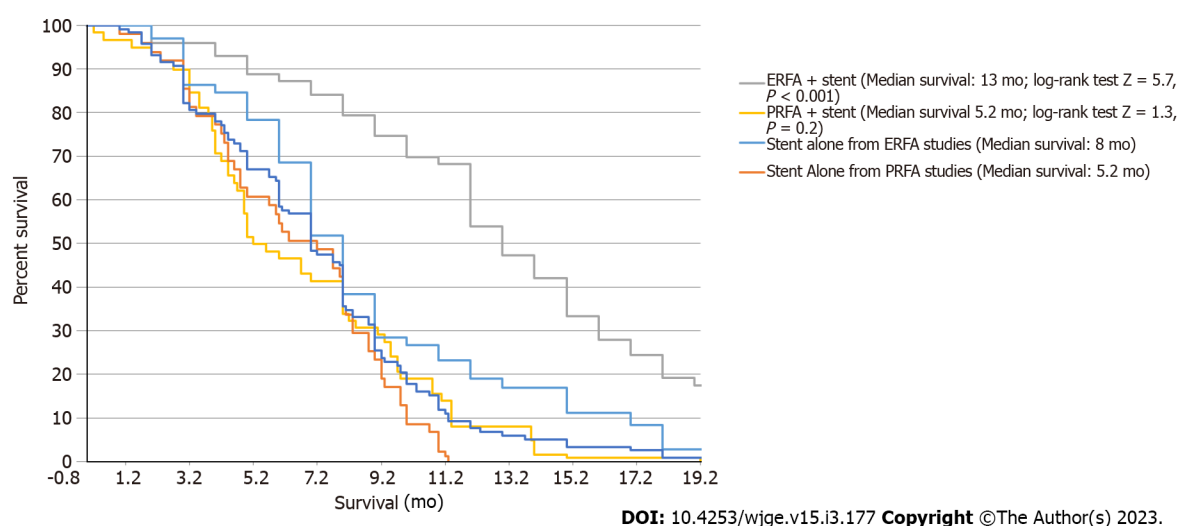
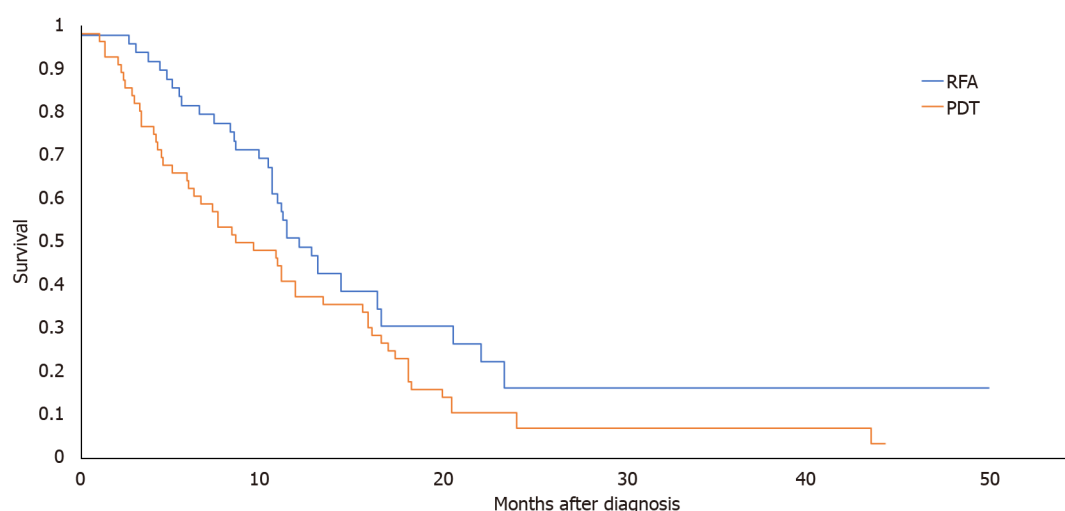


Figure 5 Kaplan-Meier survival curve of endoscopic radiofrequency ablation and percutaneous radiofrequency ablation with stenting compared to biliary stenting alone. ERFA: Endoscopic radiofrequency ablation; PRFA: Percutaneous radiofrequency ablation.

severe post-procedure pain (22.7% *vs* 4.8%, $P = 0.003$), and phototoxicity (2.7% *vs* 0%) in the pooled PDT cohort as compared to the pooled ERFA cohort (Table 6).

Quality assessment and risk for bias

The study by Strand *et al*[13] received a score of “9” out of 9 as confounders such as tumor stage, performance status, and number of procedures did not differ among cohorts. While described as a case series, the study from Schmidt and colleagues was largely retrospective and partly prospective. Designation of intervention in the prospective portion was determined by choice of the patient, thus losing a point in selection of the cohorts and receiving a score of “8” out of 9. The study performed by Wu *et al*[26] received a NOS score of “7” out of 9, as there were no cofounders corrected for. Additionally, the study by Cui *et al*[25] also received a score of “7” out of 9 because age significantly differed among study groups and was uncorrected for. The study by Bokemeyer *et al*[24] received a NOS score of “9” out of 9. In this case, confounders were adjusted for by age, extent of disease, the use of endoprotheses, and the application of systemic palliative chemotherapy. The study from Yang and colleagues was assessed using the Cochrane risk of bias tool. While subjects were randomized, patients and interventionalists could not be blinded. Additionally, there was some unclear risk for bias in this study as detailed in Supplementary Figure 1. Two studies that were published only as abstracts were not able to be assessed for bias. Detailed analysis of these scores can be seen in the appendix as Supplementary Table 3.



DOI: 10.4253/wjge.v15.i3.177 Copyright ©The Author(s) 2023.

Figure 6 Kaplan-Meier survival curve of endoscopic radiofrequency ablation with stenting compared to photodynamic therapy with stenting. RFA: Radiofrequency ablation; PDT: Photodynamic therapy.

DISCUSSION

Although it remains a relatively rare disease, the incidence of CCA continues to increase worldwide. Surgical resection remains the only curative treatment option; however, resection is only an available option in up to 30% of patients diagnosed, likely due to a variety of factors, including delayed diagnosis, which is, in large part, due to late onset of symptoms[27]. As such, for many patients, palliative approaches become the mainstay treatment options.

Our study compiles pooled data from previous investigations to better describe the roles ERFA and PRFA with stenting have in the palliation of unresectable cholangiocarcinoma and ascertain the survival benefits, thereof, while identifying adverse events that could portend poor quality of remaining life.

The meta-analytic outcomes in our study demonstrated a statistically significant improvement in both mean and median survivals when comparing ERFA to endoscopic biliary stenting alone in this cohort of patients with unresectable cholangiocarcinoma. While percutaneous RFA (PRFA) performed by capable Interventional Radiologists leads to improvement in mean survival, median survival is not impacted. While there are no studies assessing direct comparisons between ERFA and PRFA, available data does suggest superiority of ERFA with regard to median survival in these CCA patients, arguing for more widespread implementation of this palliative technique.

Safety concerns have been raised, however, given risk of stent occlusion or migration-with resulting cholangitis or delays in chemotherapy due to ensuing hyperbilirubinemia-as well as the risk of hemobilia and cholecystitis. However, the pooled data of included studies did not reveal an increase in stent occlusion rates, cholangitis, or hemobilia as compared to biliary stenting alone but did demonstrate increased risk of cholecystitis. Subgroup analyses were insufficient to conclude whether reported cholecystitis occurred in those with plastic or metallic biliary stenting. As compared to PRFA, there was an increased risk of reported cholangitis cases. However, given the lack of PRFA adverse event data reported (only one study allowed for analysis), definitive conclusions are difficult to make.

While technically feasible with reasonable safety outcomes, ERFA is an appealing option for palliation in these patients. However, the technique is limited in certain respects to degree of stricture, as severe strictures make passage of the RFA probe difficult and mild strictures may not result in adequate contact of the RFA to achieve adequate ablation. There is also a lack of consensus with regard to the timing of repeat ablation, particularly in those with successful first ablations. Further studies are needed to ascertain the optimal period between procedures as well as endoluminal and clinical parameters that would otherwise warrant repeating or avoiding the procedure.

Given the paucity of comparative studies, this meta-analysis was restricted to a small number of published studies, which could potentially overstate the benefit of the approach. Thirteen articles in our literature review were excluded in this meta-analysis due to a lack of contingency of data to separate CCA patients from those studies with other malignant biliary obstructions (ampullary and pancreatic carcinomas), and another 15 articles were excluded for inclusion of other palliative endotherapies (photodynamic therapy) or included patients in whom a previous surgical intervention was undertaken.

To this point, a recent meta-analysis by Zheng and colleagues suggested that patients undergoing ERFA for malignant biliary obstruction had a pooled survival of 9.6 mo but included all patients with malignant biliary obstruction[28]. Similarly, a separate meta-analysis compared ERFA with biliary stenting and to biliary stenting alone for malignant biliary obstructions found a mean survival of 9.4 mo

[29]. While the exact mechanism for prolonged survival is unknown, it has been postulated that the ablative process induces a systemic immune response which is then amplified by immune modulating agents resulting in improved clinical outcomes[30-32].

Our cohort of 94 cumulative patients with unresectable CCA receiving ERFA with stenting demonstrated a median survival of 13 mo. This difference may be explained by the exclusion of other etiologies for malignant biliary obstruction; technique advancement with the availability of improved cholangioscopic visualization of the malignant stricture; patient selection; or other confounders, such as stent selection.

PDT with biliary stenting is another endoscopic approach that has been well-studied as a palliative option for patients with unresectable CCA and has been shown to be superior to biliary stenting alone. While there is a paucity of studies, our meta-analysis demonstrated that in two comparative studies with available relevant contingency data, the median survival with ERFA is statistically superior than in PDT. This difference may be explained by lack of studies comparing the two modalities directly and the need for more study for adequate comparison of survival outcomes.

With lack of available studies, the direction of endoscopic palliative therapy is one that, at present, is largely center-dependent. PDT has the inconvenience of requiring two stages of intervention, one for administration of the photosensitizer and one for the delivery of therapy for tumor necrosis and cell death and also comes with the added inconvenience for the patient of avoiding direct exposure to light due to risk of skin photosensitivity. This is not the case with ERFA, which can be performed as a single procedure. It is worth noting, however, that increased rates of cholangitis and stent occlusion in ERFA cohorts would increase the need for subsequent interventions and increase costs related to repeat procedures, but this is an outcome that must also be studied further. In comparing ERFA with stenting compared to biliary stenting alone, however, there was no statistically significant difference in stent occlusion or cholangitis adverse events, so as a singular modality, safety outcomes are still comparable to biliary stenting alone while offering the benefit of longer survival as compared to biliary stenting. Interestingly, while PDT did have higher rate of stent migration, this may potentially reflect significant decrease in size of the obstructing tumor, which is a desirable outcome; this, however, was not quantified in the comparative studies and is an area for potential investigation.

CONCLUSION

In any event, endoscopic palliation of unresectable CCA with ERFA has shown significant promise in this patient population, but further studies are needed to assess our specific cohort of patients to further understand palliative, technical, and clinical outcomes, especially as they compare to other palliative therapies that extend beyond conventional biliary stenting alone.

ARTICLE HIGHLIGHTS

Research background

Further prospective studies comparing all therapeutic modalities are needed to best understand their role in the treatment of unresectable cholangiocarcinoma.

Research motivation

Endoscopic radiofrequency ablation with biliary stenting is a promising palliative therapeutic option in patients presenting with unresectable cholangiocarcinoma.

Research objectives

Endoscopic radiofrequency ablation when used in conjunction with biliary stenting showed improved survival benefit when compared to alternative palliative therapies.

Research methods

This is a comprehensive literature review of studies evaluating survival benefit and other clinical outcomes as it relates to the proposed therapeutic interventions.

Research results

To better understand, qualify, and quantify the survival outcomes of endoscopic radiofrequency ablation, percutaneous radiofrequency ablation, and photodynamic therapy in the treatment of unresectable cholangiocarcinoma as it compares to conventional therapy alone.

Research conclusions

Our motivation for this study was to better understand alternative approaches to palliative endoscopic

intervention for patients with unresectable cholangiocarcinoma.

Research perspectives

There is limited data evaluating the clinical outcomes of endoscopic radiofrequency ablation and photodynamic therapy as interventions for unresectable cholangiocarcinoma.

FOOTNOTES

Author contributions: Villa E contributed to conception and design; Rebhun J and Villa E contributed to analysis and interpretation of the data; Rebhun J, Shin CM, Villa E contributed to drafting of the article; Siddiqui UD and Villa E contributed to critical revision of the article for important intellectual content; Rebhun J, Shin CM, Siddiqui UD, Villa E contributed to final approval of the article.

Conflict-of-interest statement: Uzma Siddiqui has served as a speaker/consultant for Boston Scientific, Medtronic, and Olympus. Edward Villa has served as a consultant/speaker for ConMed as well as a speaker for Ovesco.

PRISMA 2009 Checklist statement: The authors have read the PRISMA 2009 Checklist, and the manuscript was prepared and revised according to the PRISMA 2009 Checklist.

Open-Access: This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <https://creativecommons.org/licenses/by-nc/4.0/>

Country/Territory of origin: United States

ORCID number: Jeffrey Rebhun 0000-0001-8883-009X.

S-Editor: Liu JH

L-Editor: A

P-Editor: Liu JH

REFERENCES

- Global Burden of Disease Cancer Collaboration,** Fitzmaurice C, Dicker D, Pain A, Hamavid H, Moradi-Lakeh M, MacIntyre MF, Allen C, Hansen G, Woodbrook R, Wolfe C, Hamadeh RR, Moore A, Werdecker A, Gessner BD, Te Ao B, McMahon B, Karimkhani C, Yu C, Cooke GS, Schwebel DC, Carpenter DO, Pereira DM, Nash D, Kazi DS, De Leo D, Plass D, Ukwaja KN, Thurston GD, Yun Jin K, Simard EP, Mills E, Park EK, Catalá-López F, deVeber G, Gotay C, Khan G, Hosgood HD 3rd, Santos IS, Leasher JL, Singh J, Leigh J, Jonas JB, Sanabria J, Beardsley J, Jacobsen KH, Takahashi K, Franklin RC, Ronfani L, Montico M, Naldi L, Tonelli M, Geleijnse J, Petzold M, Shrimo MG, Younis M, Yonemoto N, Breitborde N, Yip P, Pourmalek F, Lotufo PA, Esteghamati A, Hankey GJ, Ali R, Lunevicius R, Malekzadeh R, Dellavalle R, Weintraub R, Lucas R, Hay R, Rojas-Rueda D, Westerman R, Sepanlou SG, Nolte S, Patten S, Weichenthal S, Abera SF, Fereshtehnejad SM, Shiue I, Driscoll T, Vasankari T, Alsharif U, Rahimi-Movaghar V, Vlassov VV, Marcenes WS, Mekonnen W, Melaku YA, Yano Y, Artaman A, Campos I, MacLachlan J, Mueller U, Kim D, Trillini M, Eshrati B, Williams HC, Shibuya K, Dandona R, Murthy K, Cowie B, Amare AT, Antonio CA, Castañeda-Orjuela C, van Gool CH, Violante F, Oh IH, Deribe K, Soreide K, Knibbs L, Kereselidze M, Green M, Cardenas R, Roy N, Tillmann T, Li Y, Krueger H, Monasta L, Dey S, Sheikhbahaei S, Hafezi-Nejad N, Kumar GA, Sreeramareddy CT, Dandona L, Wang H, Vollset SE, Mokdad A, Salomon JA, Lozano R, Vos T, Forouzanfar M, Lopez A, Murray C, Naghavi M. The Global Burden of Cancer 2013. *JAMA Oncol* 2015; **1**: 505-527 [PMID: 26181261 DOI: 10.1001/jamaoncol.2015.0735]
- National Cancer Institute.** "Cancer stat facts: liver and intrahepatic bile duct cancer." 2020 [DOI: 10.32388/AOLRSJ]
- DeOliveira ML,** Cunningham SC, Cameron JL, Kamangar F, Winter JM, Lillemoe KD, Choti MA, Yeo CJ, Schulick RD. Cholangiocarcinoma: thirty-one-year experience with 564 patients at a single institution. *Ann Surg* 2007; **245**: 755-762 [PMID: 17457168 DOI: 10.1097/01.sla.0000251366.62632.d3]
- Esnaola NF,** Meyer JE, Karachristos A, Maranki JL, Camp ER, Denlinger CS. Evaluation and management of intrahepatic and extrahepatic cholangiocarcinoma. *Cancer* 2016; **122**: 1349-1369 [PMID: 26799932 DOI: 10.1002/cncr.29692]
- Massironi S,** Pilla L, Elvevi A, Longarini R, Rossi RE, Bidoli P, Invernizzi P. New and Emerging Systemic Therapeutic Options for Advanced Cholangiocarcinoma. *Cells* 2020; **9** [PMID: 32168869 DOI: 10.3390/cells9030688]
- Rahnamai-Azar AA,** Weisbrod AB, Dillhoff M, Schmidt C, Pawlik TM. Intrahepatic cholangiocarcinoma: current management and emerging therapies. *Expert Rev Gastroenterol Hepatol* 2017; **11**: 439-449 [PMID: 28317403 DOI: 10.1080/17474124.2017.1309290]
- Chandrasegaram MD,** Eslick GD, Mansfield CO, Liem H, Richardson M, Ahmed S, Cox MR. Endoscopic stenting versus operative gastrojejunostomy for malignant gastric outlet obstruction. *Surg Endosc* 2012; **26**: 323-329 [PMID: 21898024 DOI: 10.1007/s00464-011-1870-3]

- 8 **Sangchan A**, Kongkasame W, Pugkhem A, Jenwitheesuk K, Mairiang P. Efficacy of metal and plastic stents in unresectable complex hilar cholangiocarcinoma: a randomized controlled trial. *Gastrointest Endosc* 2012; **76**: 93-99 [PMID: [22595446](#) DOI: [10.1016/j.gie.2012.02.048](#)]
- 9 **Zoepl T**. Photodynamic therapy of cholangiocarcinoma. *HPB (Oxford)* 2008; **10**: 161-163 [PMID: [18773045](#) DOI: [10.1080/13651820801992625](#)]
- 10 **Moole H**, Tathireddy H, Dharmapuri S, Moole V, Boddireddy R, Yedama P, Uppu A, Bondalapati N, Duvvuri A. Success of photodynamic therapy in palliating patients with nonresectable cholangiocarcinoma: A systematic review and meta-analysis. *World J Gastroenterol* 2017; **23**: 1278-1288 [PMID: [28275308](#) DOI: [10.3748/wjg.v23.i7.1278](#)]
- 11 **Ortner ME**, Caca K, Berr F, Liebetruht J, Mansmann U, Huster D, Voderholzer W, Schachschal G, Mössner J, Lochs H. Successful photodynamic therapy for nonresectable cholangiocarcinoma: a randomized prospective study. *Gastroenterology* 2003; **125**: 1355-1363 [PMID: [14598251](#) DOI: [10.1016/j.gastro.2003.07.015](#)]
- 12 **Zoepl T**, Jakobs R, Arnold JC, Apel D, Riemann JF. Palliation of nonresectable bile duct cancer: improved survival after photodynamic therapy. *Am J Gastroenterol* 2005; **100**: 2426-2430 [PMID: [16279895](#) DOI: [10.1111/j.1572-0241.2005.00318.x](#)]
- 13 **Strand DS**, Cosgrove ND, Patrie JT, Cox DG, Bauer TW, Adams RB, Mann JA, Sauer BG, Shami VM, Wang AY. ERCP-directed radiofrequency ablation and photodynamic therapy are associated with comparable survival in the treatment of unresectable cholangiocarcinoma. *Gastrointest Endosc* 2014; **80**: 794-804 [PMID: [24836747](#) DOI: [10.1016/j.gie.2014.02.1030](#)]
- 14 **Gao DJ**, Hu B, Ye X, Wu J, Wang TT, Xia MX, Sun B. Radiofrequency ablation has comparable overall survival to photodynamic therapy in the treatment of cholangiocarcinoma and ampullary carcinoma. *Gastrointest Endosc* 2018; **87**: AB72 Abstract 340 [DOI: [10.1016/j.gie.2018.04.060](#)]
- 15 **Schmidt A**, Bloechinger M, Weber A, Siveke J, von Delius S, Prinz C, Schmitt W, Schmid RM, Neu B. Short-term effects and adverse events of endoscopically applied radiofrequency ablation appear to be comparable with photodynamic therapy in hilar cholangiocarcinoma. *United European Gastroenterol J* 2016; **4**: 570-579 [PMID: [27536367](#) DOI: [10.1177/2050640615621235](#)]
- 16 **Shah DR**, Green S, Elliot A, McGahan JP, Khatri VP. Current oncologic applications of radiofrequency ablation therapies. *World J Gastrointest Oncol* 2013; **5**: 71-80 [PMID: [23671734](#) DOI: [10.4251/wjgo.v5.i4.71](#)]
- 17 **Steel AW**, Postgate AJ, Khorsandi S, Nicholls J, Jiao L, Vlavianos P, Habib N, Westaby D. Endoscopically applied radiofrequency ablation appears to be safe in the treatment of malignant biliary obstruction. *Gastrointest Endosc* 2011; **73**: 149-153 [PMID: [21184881](#) DOI: [10.1016/j.gie.2010.09.031](#)]
- 18 **Wells GA**, Shea B, O'Connell D, Peterson J, Welch V, Losos M, Tugwell P. The Newcastle-Ottawa Scale (NOS) for assessing the quality of nonrandomized studies in meta-analyses." 2000 [DOI: [10.7717/peerj.14320/supp-5](#)]
- 19 **Rahmati M**, Keshvari M, Mirasuri S, Yon DK, Lee SW, Il Shin J, Smith L. The global impact of COVID-19 pandemic on the incidence of pediatric new-onset type 1 diabetes and ketoacidosis: A systematic review and meta-analysis. *J Med Virol* 2022; **94**: 5112-5127 [PMID: [35831242](#) DOI: [10.1002/jmv.27996](#)]
- 20 **Rahmati M**, Malakoutinia F. Aerobic, resistance and combined exercise training for patients with amyotrophic lateral sclerosis: a systematic review and meta-analysis. *Physiotherapy* 2021; **113**: 12-28 [PMID: [34555670](#) DOI: [10.1016/j.physio.2021.04.005](#)]
- 21 **Hu B**, Gao DJ, Zhang X, Zhang YC. 121 Endobiliary radiofrequency ablation improve overall survival of Cholangiocarcinoma: A multi-center randomized control study. *Gastrointest Endosc* 2016; **83**: AB126. [DOI: [10.1016/j.gie.2016.03.046](#)]
- 22 **Yang J**, Wang J, Zhou H, Zhou Y, Wang Y, Jin H, Lou Q, Zhang X. Efficacy and safety of endoscopic radiofrequency ablation for unresectable extrahepatic cholangiocarcinoma: a randomized trial. *Endoscopy* 2018; **50**: 751-760 [PMID: [29342492](#) DOI: [10.1055/s-0043-124870](#)]
- 23 **Sampath L**, Gardner T, Gordon SR. Tu1526 The Effect of Endoscopic Radiofrequency Ablation on Survival in Patients with Unresectable Peri-Hilar Cholangiocarcinoma. *Gastrointest Endosc* 2016; **83**: AB595 [DOI: [10.1016/j.gie.2016.03.1234](#)]
- 24 **Bokemeyer A**, Matern P, Bettenworth D, Cordes F, Nowacki TM, Heinzow H, Kabar I, Schmidt H, Ullerich H, Lenze F. Endoscopic Radiofrequency Ablation Prolongs Survival of Patients with Unresectable Hilar Cholangiocellular Carcinoma - A Case-Control Study. *Sci Rep* 2019; **9**: 13685 [PMID: [31548703](#) DOI: [10.1038/s41598-019-50132-0](#)]
- 25 **Cui W**, Wang Y, Fan W, Lu M, Zhang Y, Yao W, Li J. Comparison of intraluminal radiofrequency ablation and stents vs. stents alone in the management of malignant biliary obstruction. *Int J Hyperthermia* 2017; **33**: 853-861 [PMID: [28540797](#) DOI: [10.1080/02656736.2017.1309580](#)]
- 26 **Wu TT**, Li WM, Li HC, Ao GK, Zheng F, Lin H. Percutaneous Intraductal Radiofrequency Ablation for Extrahepatic Distal Cholangiocarcinoma: A Method for Prolonging Stent Patency and Achieving Better Functional Status and Quality of Life. *Cardiovasc Intervent Radiol* 2017; **40**: 260-269 [PMID: [27743089](#) DOI: [10.1007/s00270-016-1483-2](#)]
- 27 **Razumilava N**, Gores GJ. Cholangiocarcinoma. *Lancet* 2014; **383**: 2168-2179 [PMID: [24581682](#) DOI: [10.1016/S0140-6736\(13\)61903-0](#)]
- 28 **Zheng X**, Bo ZY, Wan W, Wu YC, Wang TT, Wu J, Gao DJ, Hu B. Endoscopic radiofrequency ablation may be preferable in the management of malignant biliary obstruction: A systematic review and meta-analysis. *J Dig Dis* 2016; **17**: 716-724 [PMID: [27768835](#) DOI: [10.1111/1751-2980.12429](#)]
- 29 **Sofi AA**, Khan MA, Das A, Sachdev M, Khuder S, Nawras A, Lee W. Radiofrequency ablation combined with biliary stent placement versus stent placement alone for malignant biliary strictures: a systematic review and meta-analysis. *Gastrointest Endosc* 2018; **87**: 944-951.e1 [PMID: [29108980](#) DOI: [10.1016/j.gie.2017.10.029](#)]
- 30 **Hansler J**, Wissniewski TT, Schuppan D, Witte A, Bernatik T, Hahn EG, Strobel D. Activation and dramatically increased cytolytic activity of tumor specific T lymphocytes after radio-frequency ablation in patients with hepatocellular carcinoma and colorectal liver metastases. *World J Gastroenterol* 2006; **12**: 3716-3721 [PMID: [16773688](#) DOI: [10.3748/wjg.v12.i23.3716](#)]
- 31 **Napoleitano C**, Taurino F, Biffoni M, De Majo A, Coscarella G, Bellati F, Rahimi H, Pauselli S, Pellicciotta I, Burchell JM,

- Gaspari LA, Ercoli L, Rossi P, Rughetti A. RFA strongly modulates the immune system and anti-tumor immune responses in metastatic liver patients. *Int J Oncol* 2008; **32**: 481-490 [PMID: [18202772](#) DOI: [10.3892/ijo.32.2.481](#)]
- 32 **Liu X**, Qin S. Immune Checkpoint Inhibitors in Hepatocellular Carcinoma: Opportunities and Challenges. *Oncologist* 2019; **24**: S3-S10 [PMID: [30819826](#) DOI: [10.1634/theoncologist.2019-IO-S1-s01](#)]



Published by **Baishideng Publishing Group Inc**
7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA

Telephone: +1-925-3991568

E-mail: bpgoffice@wjgnet.com

Help Desk: <https://www.f6publishing.com/helpdesk>

<https://www.wjgnet.com>

