

University of Groningen



Major complications and mortality after resection of intrahepatic cholangiocarcinoma

van Keulen, Anne Marleen; Büttner, Stefan; Erdmann, Joris I.; Hagendoorn, Jeroen; Hoogwater, Frederik J.H.; IJzermans, Jan N.M.; Neumann, Ulf P.; Polak, Wojciech G.; De Jonge, Jeroen; Olthof, Pim B.

Published in: Surgery (United States)

DOI: 10.1016/j.surg.2022.11.027

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version Version created as part of publication process; publisher's layout; not normally made publicly available

Publication date: 2022

Link to publication in University of Groningen/UMCG research database

Citation for published version (APA):

van Keulen, A. M., Büttner, S., Erdmann, J. I., Hagendoorn, J., Hoogwater, F. J. H., IJzermans, J. N. M., Neumann, U. P., Polak, W. G., De Jonge, J., Olthof, P. B., & Koerkamp, B. G. (Accepted/In press). Major complications and mortality after resection of intrahepatic cholangiocarcinoma: A systematic review and meta-analysis. Surgery (United States). https://doi.org/10.1016/j.surg.2022.11.027

Copyright

Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

The publication may also be distributed here under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license. More information can be found on the University of Groningen website: https://www.rug.nl/library/open-access/self-archiving-pure/taverneamendment.

Take-down policy If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Downloaded from the University of Groningen/UMCG research database (Pure): http://www.rug.nl/research/portal. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.

Surgery xxx (2022) 1-10



Contents lists available at ScienceDirect

Surgery



journal homepage: www.elsevier.com/locate/surg

Major complications and mortality after resection of intrahepatic cholangiocarcinoma: A systematic review and meta-analysis

Anne-Marleen van Keulen, MD^a, Stefan Büttner, MD, PhD^a, Joris I. Erdmann, MD, PhD^b, Jeroen Hagendoorn, MD, PhD^c, Frederik J.H. Hoogwater, MD, PhD^d, Jan N.M. IJzermans, MD, PhD^a, Ulf P. Neumann, MD, PhD^e, Wojciech G. Polak, MD, PhD^a, Jeroen De Jonge, MD, PhD^a, Pim B. Olthof, MD, PhD^a, Bas Groot Koerkamp, MD, PhD^{a,*}

^a Department of Surgery, Erasmus MC Cancer Institute, Rotterdam, the Netherlands

^b Department of Surgery, Amsterdam University Medical Center, the Netherlands

^c Department of Surgery, Regional Academic Cancer Center Utrecht, the Netherlands

^d Department of Surgery, section Hepato-Pancreato-Biliary Surgery and Liver Transplantation, University of Groningen, University Medical Center

Groningen, the Netherlands

^e Department of Surgery, Maastricht University Medical Center, the Netherlands

ARTICLE INFO

Article history: Accepted 20 November 2022 Available online xxx

ABSTRACT

Background: Evaluation of morbidity and mortality after hepatic resection often lacks stratification by extent of resection or diagnosis. Although a liver resection for different indications may have technical similarities, postoperative outcomes differ. The aim of this systematic review and meta-analysis was to determine the risk of major complications and mortality after resection of intrahepatic cholangiocarcinoma. *Methods:* Meta-analysis was performed to assess postoperative mortality (in-hospital, 30-, and 90-day) and major complications (Clavien-Dindo grade \geq III). *Results:* A total of 32 studies that reported on 19,503 patients were included. Pooled in-hospital, 30-day, and 90-day mortality were 5.9% (95% confidence interval 4.1–8.4); 4.6% (95% confidence interval 4.0–5.2); and 6.1% (95% confidence interval 5.0–7.3), respectively. Pooled proportion of major complications was 22.2% (95% confidence interval 1.7–27.5) for all resections. The pooled 90-day mortality was 3.1% (95% confidence interval 4.9–8.4); 4.9% confidence interval 6.9–18.7) for extended resections (*P* =.001). Major complications were 38.8%

(95% confidence interval 29.5–49) after a major hepatectomy compared to 11.3% (95% confidence interval 5.0 –24.0) after a minor hepatectomy (P = .001). Asian studies had a pooled 90-day mortality of 4.4% (95% confidence interval 3.3–5.9) compared to 6.8% (95% confidence interval 5.6–8.2) for Western studies (P = .02). Cohorts with patients included before 2000 had a pooled 90-day mortality of 5.9% (95% confidence interval 4.8–7.3) compared to 6.8% (95% confidence interval 5.1–9.1) after 2000 (P = .44).

Conclusion: When informing patients or comparing outcomes across hospitals, postoperative mortality rates after liver resection should be reported for 90-days with consideration of the diagnosis and the extent of liver resection.

© 2022 The Author(s). Published by Elsevier Inc. This is an open access article under the CC BY license (http://creativecommons.org/licenses/by/4.0/).

Introduction

Cholangiocarcinoma is a heterogenous group of malignancies in the biliary tree and accounts for 3% of all gastrointestinal cancers worldwide.^{1,2} Intrahepatic cholangiocarcinoma (iCCA)

* Reprint requests: Bas Groot Koerkamp, MD, PhD, Department of Surgery, Erasmus MC Cancer Institute, Doctor Molewaterplein 40, 3015 GD Rotterdam, the Netherlands.

arises from the epithelial cells of the peripheral bile ducts proximal to the second-order bile ducts.³ Intrahepatic cholangiocarcinoma represents around 20% of all cholangiocarcinoma and its incidence in Western countries is 1 to 2 per 100,000.^{4,5} The incidence of cholangiocarcinoma in Asian countries exceeds that in Western countries, up to >6 per 100,000.⁶ Treatment and diagnosis of patients with iCCA entails many challenges. Patients with iCCA are initially asymptomatic and 20% of patients are diagnosed in the absence of symptoms.⁷ When symptomatic, most patients present with unresectable disease due to locally advanced or metastatic disease.

https://doi.org/10.1016/j.surg.2022.11.027

0039-6060/© 2022 The Author(s). Published by Elsevier Inc. This is an open access article under the CC BY license (http://creativecommons.org/licenses/by/4.0/).

E-mail address: b.grootkoerkamp@erasmusmc.nl (B.G. Koerkamp); Twitter: @BasGrootKoerkam

ARTICLE IN PRESS

A.-M. van Keulen et al. / Surgery xxx (2022) 1–10

About 20% of patients are eligible for complete surgical resection.⁸ Five-year overall survival (OS) rate after curative-intent resection of iCCA is about 30% and median OS about 30 months after surgery.^{9–11} Resection usually involves a major hepatectomy (82%) (3 segments or more) and sometimes resection of the (common) bile duct (23%) requiring an additional hepatico-jejunostomy.¹² In addition, most patients have underlying liver disease or postcholestatic liver dysfunction. Therefore, these extensive resections come with substantially higher risks than for instance resections for colorectal metastasis. The most frequent postoperative complications are liver failure, biliary leakage, and intraabdominal abscess.^{11,13,14} Postoperative morbidity and mortality rates vary in the literature. Postoperative major complication risks vary from 18% to 52%,^{15,16} whereas overall 90-day postoperative mortality ranges from 2% to 11%.^{16,17}

The evaluation of morbidity and mortality after hepatic resection often lacks stratification by the extent of resection or diagnosis. Although a hepatectomy for different indications may have technical similarities, postoperative outcomes differ.¹⁸ Due to the low incidence of iCCA, postoperative morbidity and mortality rates are mainly derived from observational cohort studies or case series. Therefore, the aim of this systematic review and meta-analysis was to determine the risk of postoperative major complications and mortality after resection of iCCA. With subgroup analyses for the extent of liver resection, region, and time period.

Methods

This systematic review and meta-analysis was performed in accordance with the Preferred Reporting Items for Systematic Review and Meta-Analyses statement. 19

Literature search

A librarian was consulted to perform a systematic search on February 19, 2021. Databases of MEDLINE, Embase, the Cochrane Library, and Web of Science were queried. A combination of keywords and Medical Subject Headings terms were used for the search: bile duct neoplasms, intrahepatic cholangiocarcinoma, hepatectomy, postoperative morbidity, postoperative mortality and variations thereof. No restriction on date, language, or publication type was applied in the search. The complete search strategy is presented in Supplementary Table S1. Two independent authors (A.K. and S.B.) screened the title and abstract of each identified publication for eligibility. Publications that seemed eligible for inclusion were retrieved for full text reading by A.K. Discrepancies at any stage were resolved by a third author (P.B.O.).

Eligibility criteria

All studies that reported on complications and mortality after resection of iCCA were eligible for inclusion. Non-English and non-original publications such as reviews, case reports, letters, editorials, or conference abstracts were excluded. Studies including patients who underwent a resection before 1990 were excluded. Studies that reported a combined study population (eg, intrahepatic- and extrahepatic cholangiocarcinoma) were only included if outcomes were presented separately for iCCA. If a population had mixed pathology (eg, iCCA and hepatocellular carcinoma), the study was excluded. Studies that lacked data on the outcomes of interest were excluded, as well as studies that reported on treatments other than initial complete resection of iCCA (eg, associating liver partition and portal vein ligation for staged hepatectomy and resection of recurrences). In case of overlapping cohorts, the largest or the most recent series was included. Studies with <50 patients were excluded.

Data collection

The extracted data included study characteristics (ie. author. vear of publication, inclusion period, and number of patients). patient characteristics (ie, age, sex, and American Society of Anesthesiologists classification [ASA]), preoperative characteristics (ie, portal vein embolization), operative characteristics (ie, type of resection, vascular reconstruction, and biliary reconstruction), and postoperative details (ie, major complications and mortality). Authors from some studies were contacted for lacking data. The primary outcome was postoperative mortality, defined as in-hospital, 30-day, or 90-day mortality. The secondary outcome was major complications (Clavien-Dindo grade ≥III). Subgroup analyses were prespecified for 90-day mortality and major complications in relation to the extent of liver resection (ie, minor, major, and extended), period (ie, start of inclusion period before or after 2000), and region (ie, Asian or Western studies). A study was classified as Asian or Western if the origin of the participating centers originated for at least 80% from Asian or Western countries. A minor hepatectomy was defined as resection of <3 Couinaud segments, a major hepatectomy was defined as resection of >3 Couinaud segments, and an extended hepatectomy was defined as resection of >5 Couinaud segments.

Quality assessment

Risk of bias was assessed by the Joanna Briggs Institute checklist that is specifically designed for case series.²⁰ The checklist consists of 10 predefined items that can be specified to questions particularly relevant for the interest of this systematic review, which can be answered with 'yes,' 'no,' and 'unclear or not applicable' (Supplementary Table S2).

Statistical analysis

The categorical values are presented as numbers and percentages. Continuous data are presented as mean with SD. Studies were pooled separately for in-hospital, 30-day, and 90-day mortality. Studies were pooled for postoperative morbidity if they reported major complications (Clavien-Dindo grade \geq III). The I^2 statistic was used to quantify the heterogeneity across studies. An I^2 value >50%indicated significant heterogeneity. A random-effects model was used for all analyses. Pooled analyses were visualized with forest plots. Statistical analyses were performed using R version 4.1.1 (http://www.r-project.org; R Foundation for Statistical Computing, Vienna, Austria).

Results

The search and screening process of the literature search is presented in Figure 1. A total of 9,821 records were identified through database searching. The duplicate records were removed. The title and abstract of 7,076 records were screened for eligibility. This led to full-text assessment of 184 studies, of which 32 were included in the meta-analysis.

Study characteristics

The 32 included studies reported on 19,503 patients who underwent resection of iCCA between 1990 and 2018. The main characteristics of the included studies are presented in Table I. Fourteen studies had a multicenter design, 11,15,21-32 of which 4

A.-M. van Keulen et al. / Surgery xxx (2022) 1-10

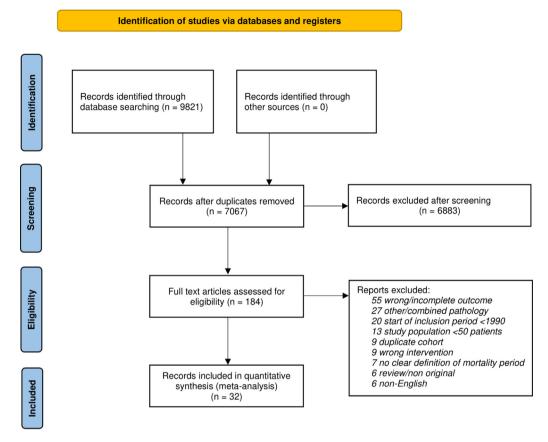


Figure 1. Preferred Reporting Items for Systematic Review and Meta-Analyses flowchart of the study selection process.

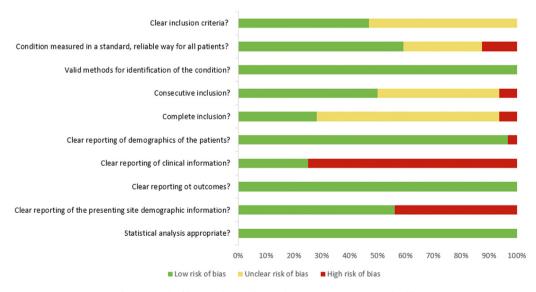


Figure 2. Risk of bias graph according to the Joanna Briggs Institute checklist.

studies queried nationwide databases.^{29–32} Of the 14 multicenter studies, 5 studies were multinational.^{15,24–27} None of these studies presented the results separately per center or per country. Twenty-two studies originated from Western countries.^{11,15,16,18,21,22–39} whereas 10 studies originated from Asian countries.^{13,22,40–47} Western countries included the USA,^{29–31} Germany,^{23,32–35,38,39} France,^{16,36} Italy,^{11,21,37} Spain,¹⁸ and the Netherlands.²⁸ Asian countries included Japan,^{22,43,45} China,^{13,41,42,44,46,47} and the Republic of Korea.⁴⁰ All studies had a retrospective design.

Patient characteristics

Twenty-nine studies (representing 16,199 patients) had a pooled minor hepatectomy proportion of 25.9% (95% CI 18-35.7).^{11,13,15,16,18,21–29,31–35,37–43,45–47} Thirty studies (representing 16,306 patients) had a pooled major hepatectomy (including extended hepatectomies) proportion of 72.8% (95% CI 62.5-81.1).^{11,13,15,16,18,21–29,31–43,45–47} As a subgroup of major hepatectomies, 22 studies (representing 13,007 patients) had a pooled

A.-M. van Keulen et al. / Surgery xxx (2022) 1–10

Table 1 Study characteristics

Author Country Inclusion Liver Male, n (%) Median ASA 3-4 n (%) Resection type Bile duct n (%) Vascular PVE n (%) period resections age (y) (LH/RH/ELH/ resections n (%) ERH) Asian studies An⁴⁷ 2004-2013 14/10/1/1 China 66 (58) 114 56 Cho⁴⁰ Korea 2001-2007 63 41 (65) 61.4 8/21/13/5 3 (5) Li⁴¹ China 2001-2009 144 91 (63) 72/16/25/9 Luo⁴² China 2007-2011 1281 912 (68) Ma¹³ 1991-2013 61 11/13/9/21 V: 4(4)China 107 59(55)15(14)Miyata⁴³ Japan 2002-2016 60 14 (23) 5(8) Morimoto²² Japan 1991-2000 51 27 (53) 13/6/7/8 16 (31) V: 2 (4) Si⁴⁴ China 2006-2010 702 428 (61) Yoh⁴⁵ 1993-2014 49 (34) Japan 144 85 (59) _ Zhu⁴⁶ 28/17/0/0 China 2012-2017 51 (61) 83 _ 8(10)Western studies 2008-2018 Bartsch³³ Germany 150 73 (49) 64.2 86 (57) 10/25/22/26 A: 0 (0), V: 18 (12) 1996-2018 51 (19) Beetz³ Germany 269 134 (50) 62 85/41/31/64 Bektas³⁵ Germany 1996-2010 158 84 (53) 61 2(1)Bergeat³⁶ France 1997-2013 107 82 (77) 10(9) A: 3 (3), V: 7 (7) 14(13) Buettner⁴ Multinational 1990-2017 1013 540 (55) 59 348 (34) Conci²¹ 1995-2015 270 137 (51) 68 V: 15 (6) Italv 984/827/-/-Filmann³² 2010-2015 Germany 4667 Guglielmi³⁷ Italy 1990-2007 52 32 (62) 17/9/2/5 V: 2 (4) 2000-2016 57 (50) Hobeika¹⁰ France 115 19 (17) 11(10)lutric³ USA 1998-2011 881 392 (45) 140/140/39/51 USA 2004-2014 1046 (46) Lee² 2256 Liu³⁰ USA 2005-2012 2089 1436 (69) _ Lurje³⁸ Germany 2011-2016 34 (48) 42 (59) 8(11) 71 Merath²⁷ Multinational 1993-2015 687 370 (54) 61 268 (39) Nickkholgh³⁹ 2001-2015 45/41/21/34 Germany 190 107(56)63 4(2)Olthof⁵⁰ Netherlands 2014-2017 97 53 (55) 67 26 (27) 24/12/5/13 13 (13) Rafecas¹⁷ Spain 1996-2017 67 45 (67) 66 20/11/4/4 4(6) Reames²⁶ Multinational 1990-2016 1087 594 (55) 438 (40) 218/179/107/140 190 (18) V: 98 (9) Ribero¹¹ 1990-2008 434 243 (56) 65 V: 14(3) Italy -/-/19/65 84 (19) Schnitzbauer²³ 250 (51) Germany 2004-2013 488 67 Spolverato²⁵ Multinational 1990-2013 583 302 (52) 59.9 Zhang¹⁵ Multinational 1990-2016 1023 569 (56) 59 202/161/99/128 177(17)

A, hepatic artery reconstruction; ASA, American Society of Anesthesiologists; ELH, extended left hemihepatectomy; ERH, extended right hemihepatectomy; LH, left hemihepatectomy; PVE, portal vein embolization; RH, right hemihepatectomy; V, portal vein reconstruction.

extended hepatectomy proportion of 22.8% (95% CI 17.3–29.4).^{11,13,15,18,22,26,28,29,31–42,45,47} Six studies (representing 647 patients) had a pooled rate of preoperative portal vein embolization of 8.2% (95% CI 5.0–13.1) (Supplementary Figure S1, *A*).^{16,18,28,36,38,39} The pooled rate of a vascular resection was 10.2% (95% CI 6.7–15.3) and involved mostly a portal vein reconstruction (Supplementary Figure S1, *B*).^{13,15,16,21,26,28,33–36,39,45}

Critical appraisal and risk of bias

The individual score of risk-of-bias per study is given in Supplementary Table S3. The overall risk of bias for each item on the Joanna Briggs Institute checklist is presented in Figure 2. Incomplete reporting of clinical information (ASA classification, extent of resection) occurred in 75% (24/32) of the studies. Incomplete reporting of the demographic information occurred in 43.8% (14/32) of the studies. No multicenter studies presented the individual results of each center. Follow-up data were unclear or incomplete in 71.9% (23/32) of the studies.

Postoperative mortality

Supplementary Table S4 presents the mortality rates per study. In-hospital mortality was reported by 8 studies (representing 7,639 patients) and ranged from 1.6% (1/63) to 11% (513/4,667) across studies.^{22,23,30,32,35,37,38,40} The pooled in-hospital mortality was 5.9% (95% CI 4.1–8.4) (Figure 3, A). Eleven studies (representing 6,847 patients) described 30-day mortality rates ranging from 0% (0/83) to 8% (12/150) across studies.^{13,15,26,27,29,31,33,34,39,46,47} The pooled 30-day mortality was 4.6% (95% CI 4.0–5.2) (Figure 3, *B*). Sixteen studies (representing 8,607 patients) reported on 90-day mortality ranging from 1.5% (1/67) to 11.3% (13/115) across studies.^{11,13,15,16,18,21,23–26,29,33,34,36,41,43,44} The pooled 90-day mortality of these studies was 6.1% (95% CI 5.0–7.3; Figure 3, *C*). Six studies reported both on 30- and 90-day mortality.^{13,15,26,29,33,34} The pooled difference between 30- and 90-day mortality was 2.5% (95% CI 1.6–3.8) (Supplementary Figure S2, *A*). Six studies (representing 16 patients) reported the proportion in which liver failure was the cause of mortality, for which the pooled proportion was 29.1% (95% CI 18.7–42) (Supplementary Figure S2, *B*).^{22,33–36,47}

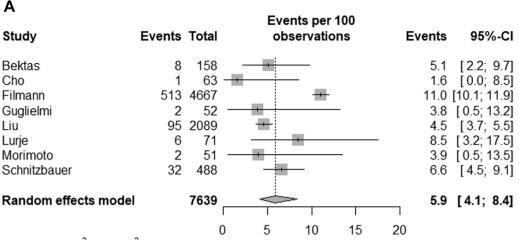
Major complications

Twenty studies (representing 8,213 patients) reported on major complications after resection of iCCA, ranging from 4% (51/1,281) to 52.2 (60/115) across studies.^{11,13,15,16,21,23-26,28,33-38,42-45} The pooled major complication rate after resection of iCCA (Clavien-Dindo grade \geq III) was 22.2% (95% CI 17.7–27.5; Figure 3, *D*).

Major complications and mortality in relation to the extent of resection

Ninety-day mortality rates were reported for minor resections in 2 studies (representing 452 patients),^{15,33} major resections in 4

A.-M. van Keulen et al. / Surgery xxx (2022) 1-10



Heterogeneity: $l^2 = 92\%$, $\tau^2 = 0.1470$, p < 0.01

D

В			Events per 100		
Study	Events	Total	observations	Events 95%	-CI
An	1	114	-	0.9 [0.0; 4	
Bartsch	12	150	<u>.</u>	8.0 [4.2; 13	
Beetz	11	269		4.1 [2.1; 7	[.2]
Jutric	52	881	÷ •	5.9 [4.4; 7	[.7]
Lee	88	2256	-	3.9 [3.1; 4	1.8]
Ma	3	107		2.8 [0.6; 8	3.0]
Merath	30	687		4.4 [3.0; 6	5.2]
Nickkholgh	13	190		6.8 [3.7; 11	.4]
Reames	52	1087	- 	4.8 [3.6; 6	5.2]
Zhang	49	1023	- <u>+</u> -	4.8 [3.6; 6	3.3]
Zhu	0	83		0.0 [0.0; 4	1.3]
Random effects model		6847	÷ 	4.6 [4.0; 5	.2]
		() 5 10 15	20	
Heterogeneity: I^2 = 36%, τ	² = 0.0118	, p = 0.			

Figure 3. Forest plots of postoperative mortality and morbidity rates after resection of intrahepatic cholangiocarcinoma. (A) in-hospital mortality, (B) 30-day mortality, (C) 90-day mortality, and (D) major complications (Clavien-Dindo grade \geq III) after resection of intrahepatic cholangiocarcinoma.

studies (representing 892 patients),^{15,33,36,41} and extended resections in 2 studies (representing 114 patients).^{33,36} The subgroup analysis revealed a pooled 90-day mortality of 3.1% (95% Cl 1.8–5.2) for a minor hepatectomy, 7.4% (95% Cl 5.9–9.3) for all major hepatectomies, and 11.4% (95% Cl 6.7–18.7) after an extended hepatectomy (P = .001) (Figure 4, A).

Six studies (representing 932 patients) reported on major complications specifically after a major hepatectomy, which ranged from 25.6% (10/39) to 59.3% (16/27) across studies.^{15,16,28,33,36,43} Two studies (representing 452 patients) reported on major complications after a minor hepatectomy, which ranged from 7.2% (30/415) to 21.6% (8/37) across studies.^{15,33} The pooled major complication rate after a major hepatectomy was 38.8% (95% Cl 29.5–49) compared to 11.3% (95% Cl 5.0–24) after a minor hepatectomy (P = .001; Figure 4, B).

Mortality in relation to Asian versus Western studies

Ten studies (representing 2,749 patients) originated from Asian centers and 22 studies (representing 16,754 patients) originated

from Western centers. The pooled in-hospital mortality was 2.6% (95% CI 0.9–7.8) in Asian studies and 6.5% (95% CI 4.1–8.4) in Western studies (P = .12; Supplementary Figure S3, A). The pooled 30-day mortality was 1.3% (95% CI 0.4–4.3) in Asian studies and 4.8% (95% CI 4.2–5.5) in Western studies (P = .03; Supplementary Figure S3, B). The pooled 90-day mortality was 4.4% (95% CI 3.3–5.9) in Asian studies and 6.8% (95% CI 5.6–8.2) in Western studies (P = .02; Figure 4, C).

Mortality in relation to inclusion period

Ninety-day mortality risks were reported by 9 studies (representing 4,690 patients) that started the inclusion period before the year 2000,^{11,13,15,24–26,34,36,39} and 7 studies (representing 3,917 patients) that started the inclusion period after the year 2000.^{16,23,29,33,41,43,44} The pooled 90-day mortality was 6.8% (95% CI 5.1–9.1) for studies that started the inclusion period after 2000, and 5.9% (95% CI 4.8–7.3) for studies that started the inclusion period before 2000 (P = .44; Figure 4, D).

A.-M. van Keulen et al. / Surgery xxx (2022) 1–10

C Study	Events	Total	Events per 100 observations	Events	95%-CI
Bartsch	13	150		8.7	[4.7; 14.4]
Beetz	21	269		7.8	[4.9; 11.7]
Bergeat	10	107		9.3	[4.6; 16.5]
Buettner	61	1013		6.0	[4.6; 7.7]
Conci	9	270		3.3	[1.5; 6.2]
Hobeika	13	115		11.3	[6.2; 18.6]
Lee	167	2256		7.4	[6.4; 8.6]
Li	5	144		3.5	[1.1; 7.9]
Ма	7	107		6.5	[2.7; 13.0]
Miyata	2	62		3.2	[0.4; 11.2]
Rafecas	1	67		1.5	[0.0; 8.0]
Reames	88	1087		8.1	[6.5; 9.9]
Ribero	23	434		5.3	[3.4; 7.8]
Schnitzbauer	49	488		10.0	[7.5; 13.1]
Si	31	702		4.4	[3.0; 6.2]
Spolverato	18	583		3.1	[1.8; 4.8]
Zhang	60	1023		5.9	[4.5; 7.5]
Random effects mode	I	8877		6.1	[5.0; 7.3]
		(0 5 10 15	20	
Heterogeneity: $I^2 = 69\%$	$^{2} = 0.1050$	n < 0			

Heterogeneity: $I^2 = 69\%$, $\tau^2 = 0.1050$, p < 0.01

D			Evente per 100	
Study	Events	Total	Events per 100 observations	Events 95%-CI
Bartsch	52	150	i —	34.7 [27.1; 42.9]
Beetz	94	269	_ _	34.9 [29.3; 41.0]
Bektas	33	158		20.9 [14.8; 28.1]
Bergeat	28	107		26.2 [18.1; 35.6]
Buettner	186	1013	-	18.4 [16.0; 20.9]
Conci	48	270		17.8 [13.4; 22.9]
Guglielmi	10	52		19.2 [9.6; 32.5]
Hobeika	60	115		52.2 [42.7; 61.6]
Luo	51	1281	+	4.0 [3.0; 5.2]
Lurje	28	71		39.4 [28.0; 51.7]
Ma	30	107		28.0 [19.8; 37.5]
Miyata	15	62		24.2 [14.2; 36.7]
Olthof	25	97		25.8 [17.4; 35.7]
Reames	207	1087		19.0 [16.7; 21.5]
Ribero	89	434		20.5 [16.8; 24.6]
Schnitzbauer	119	488		24.4 [20.6; 28.4]
Si	86	702	-	12.3 [9.9; 14.9]
Spolverato	106	583		18.2 [15.1; 21.6]
Yoh	36	144		25.0 [18.2; 32.9]
Zhang	186	1023	-	18.2 [15.9; 20.7]
Random effects mode	I	8213		22.2 [17.7; 27.5]
2	2	(0 10 20 30 40 50 6	0

0 Heterogeneity: I^2 = 94%, τ^2 = 0.3846, ρ < 0.01

Figure 3. (continued).

A.-M. van Keulen et al. / Surgery xxx (2022) 1–10

A Study	Events Total	Events per 100 observations	Events 95%-Cl
Hepatectomy = Extend Bartsch Bergeat Random effects mode Heterogeneity: $I^2 = 0\%$, τ^2	11 87 2 27 - ! 114		12.6 [6.5; 21.5] 7.4 [0.9; 24.3] 11.4 [6.7; 18.7]
Hepatectomy = All May Bartsch Bergeat Li Zhang Random effects mode Heterogeneity: / ² = 37%,	12 113 2 27 - 5 144 - 47 608 92	*	 10.6 [5.6; 17.8] 7.4 [0.9; 24.3] 3.5 [1.1; 7.9] 7.7 [5.7; 10.1] 7.4 [5.9; 9.3]
Hepatectomy = Minor Bartsch Zhang Random effects mode Heterogeneity: $I^2 = 0\%$, τ^2	$p^2 = 0, p = 0.885$	■	2.7 [0.1; 14.2] 3.1 [1.7; 5.3] 3.1 [1.8; 5.2]
Random effects mode Heterogeneity: $l^2 = 64\%$,	0	5 10 15 20 7	6.2 [4.1; 9.3]

Test for subgroup differences: $\chi_2^2 = 13.45$, df = 2 (p = 0.001)

B Study	Events	Total	Events per 100 observations	Events	95%-CI	
Hepatectomy = Minor Bartsch Zhang	8 30	37 415	* * ·	21.6 7.2	[9.8; 38.2] [4.9; 10.2]	
Random effects model Heterogeneity: $I^2 = 88\%$, τ^2		452 , p = 0.0	004		[5.0; 23.5]	
Hepatectomy = Major						
Bartsch	44	113		38.9	[29.9; 48.6]	
Bergeat	16	27			[38.8; 77.6]	
Hobeika	47	91			[40.9; 62.3]	
Miyata	10	39			[13.0; 42.1]	
Olthof	22	54		40.7	[27.6; 55.0]	
Zhang	156	608			[22.2; 29.3]	
Random effects model		932		38.8	[29.5; 48.9]	
Heterogeneity: $I^2 = 87\%$, τ^2	² = 0.1979	, p < 0.	001			
Random effects model		1384		30.6	[19.4; 44.8]	
		C		80		
Heterogeneity: $I^2 = 94\%$, $\tau^2 = 0.6924$, $p < 0.001$						
Test for subgroup difference	$ac \cdot x^2 - 1$	0 11 dt	(-1)(n - 0.001)			

Test for subgroup differences: $\chi_1^2 = 10.44$, df = 1 (p = 0.001)

Figure 4. Forest plots of subgroup analyses. (A) 90-day mortality after a minor, major, or an extended hepatectomy (subgroup of major); (B) major complications (Clavien-Dindo grade \geq III) after a minor and major hepatectomy; (C) 90-day mortality in Asian studies and Western studies; and (D) 90-day mortality for studies that started their inclusion period before or after the year 2000.

ARTICLE IN PRESS

A.-M. van Keulen et al. / Surgery xxx (2022) 1–10

С		Events per 100	
Study	Events Total	observations	Events 95%-CI
Center = Asian		1	
Li	5 144		3.5 [1.1; 7.9]
Ma	7 107		6.5 [2.7; 13.0]
Miyata	2 62		3.2 [0.4; 11.2]
Si	31 702	-	4.4 [3.0; 6.2]
Random effects mode	1015	\diamond	4.4 [3.3; 5.9]
Heterogeneity: $I^2 = 0\%$, τ^2			•
Center = Western			
Bartsch	13 150		8.7 [4.7; 14.4]
Beetz	21 269		7.8 [4.9; 11.7]
Bergeat	10 107		→ 9.3 [4.6; 16.5]
Buettner	61 1013	- <u></u>	6.0 [4.6; 7.7]
Hobeika	13 115	-	→ 11.3 [6.2; 18.6]
Lee	167 2256		7.4 [6.4; 8.6]
Rafecas	1 67		1.5 [0.0; 8.0]
Reames	88 1087		8.1 [6.5; 9.9]
Ribero	23 434		5.3 [3.4; 7.8]
Schnitzbauer	49 488		10.0 [7.5; 13.1]
Spolverato	18 583		3.1 [1.8; 4.8]
Zhang	60 1023		5.9 [4.5; 7.5]
Random effects mode			6.8 [5.6; 8.2]
Heterogeneity: $I^2 = 69\%$,	$t^2 = 0.0827, p < 0.$	01	
Random effects mode	I 8607		6.3 [5.2; 7.5]
		1 1 1	
1 later approximation $l^2 = 0.70$			5

Heterogeneity: $I^2 = 67\%$, $\tau^2 = 0.0916$, p < 0.01Test for subgroup differences: $\chi_1^2 = 5.88$, df = 1 (p = 0.02)

D			Events per 100)		
Study	Events	Total	observations	Event	s 95%-Cl	
Inclusion = Before 200	0					
Beetz	21	269		7	8 [4.9; 11.7]	
Bergeat	10	107			3 [4.6; 16.5]	
Buettner	61	1013			0 [4.6; 7.7]	
Ма	7	107		6	5 [2.7; 13.0]	
Rafecas	1	67 -			5 [0.0; 8.0]	
Reames	88	1087		8	1 [6.5; 9.9]	
Ribero	23	434		5	.3 [3.4; 7.8]	
Spolverato	18	583	—	3	1 [1.8; 4.8]	
Zhang		1023			.9 [4.5; 7.5]	
Random effects mode		4690		5	.9 [4.8; 7.3]	
Heterogeneity: I ² = 63%, τ	² = 0.0635	, p < 0.0)1			
Inclusion = After 2000			_			
Bartsch	13	150			.7 [4.7; 14.4]	
Hobeika	13	115	-		.3 [6.2; 18.6]	
Lee		2256			4 [6.4; 8.6]	
Li	5	144			.5 [1.1; 7.9]	
Miyata	2	62			2 [0.4; 11.2]	
Schnitzbauer	49	488			.0 [7.5; 13.1]	
Si Bandam offecto model	31	702 3917			4 [3.0; 6.2]	
Random effects mode				0.	.8 [5.1; 9.1]	
Heterogeneity: $I^2 = 72\%$, τ	= 0.1103	, p < 0.0)1			
Random effects model		8607		6	.3 [5.2; 7.5]	
		ſ			- / -	
		0	5 10 1	15 20		
Heterogeneity: $I^2 = 67\%$, τ	² = 0.0916	p < 0.0)1			
Test for subgroup differences: $\chi^2 = 0.59$ df = 1 ($\rho = 0.44$)						

Test for subgroup differences: $\chi_1^2 = 0.59$, df = 1 (p = 0.44)

Figure 4. (continued).

Discussion

This systematic review and meta-analysis on postoperative major complications and mortality after resection of iCCA included 32 studies that reported on 19,503 patients. The pooled in-hospital mortality was 5.9%, the 30-day mortality 4.6%, and the 90-day mortality 6.1%. The major complication (Clavien-Dindo grade \geq III) risk after resection of iCCA was 22.2%. Differences were found for 90-day mortality was 3.1% for minor resections, 7.4% for major resections, and 11.4% for extended resections (P = .001). The major complication rate was higher after a major hepatectomy when compared to a minor hepatectomy (38.8% vs 11.3%; P = .001). The pooled 90-day mortality was lower for Asian studies when compared to Western studies (4.4% vs 6.8%; P = .02). Studies with patients included before 2000 had a similar pooled 90-day mortality as after 2000 (5.9% vs 6.8%; P = .44).

Liver resections for different indications are associated with different mortality rates, despite technical similarities.²⁸ Shubert et al suggested that outcomes after hepatectomy should be stratified by the diagnosis, by showing differences in mortality for different indications using National Surgical Quality Improvement Program data of >7,000 patients.¹⁸ The highest 30-day mortality rates were observed for cholangiocarcinoma (8.2% for intrahepatic and perihilar combined) and hepatocellular carcinoma (5.2%), which were classified as 'high risk' diagnoses. These were followed by lower risk diagnoses such as metastatic disease (1.3%), gallbladder cancer (1%), and benign neoplasms (0.5%). This study could not be included in the current review because mortality was not reported for intrahepatic cholangiocarcinoma alone. A large cohort of >100,000 liver resections in Germany for all indications between 2010 and 2015 also found substantial difference in mortality rates depending on the diagnosis.³² In-hospital mortality was 5.5% for colorectal liver metastases, 7.1% for gallbladder cancer, 9.3% for hepatocellular carcinoma, and 11% for iCCA. A fair comparison of postoperative mortality after liver resection requires consideration of diagnosis.

Mortality outcomes should not only be stratified according to diagnosis, but also by the extent of the resection. In the same German cohort, in-hospital mortality (including liver resections for all diagnoses) was 3.8% for a segmental resection, 9.1% for a major hepatectomy, and 16.2% for an extended hepatectomy. The inhospital mortality of an extended hepatectomy for iCCA (n = 709) was 21.8%, which is higher than the 11.4% in the pooled analysis for 90-day mortality in the present study.³² A study on >1,000 liver resections for iCCA found that 30-day mortality was 5 times higher and 90-day mortality was 2.5 times higher after a major compared to a minor liver resection.¹⁵

Thirty-day mortality has been used as a benchmark to assess the quality of major surgical procedures. In the present study, the pooled difference between 30-day and 90-day mortality was 2.5%, indicating that surgery related death may be underestimated at 30 days, postoperatively. Posthepatectomy liver failure (PHLF) is a major cause of death after liver resection, and 25% of patients die from PHLF >30 days after resection.⁴⁸ Posthepatectomy liver failure requires complex treatment at the intensive care unit, which prolongs hospital stay and may result in slow physical deconditioning and death eventually. Truant et al found a median time to postoperative mortality of 31 days for patients (ie, 84.6%) died from PHLF, often after >30 days.⁴⁹ Therefore, 90-day postoperative mortality is a better outcome measure when reporting outcomes of liver resection.

A difference was found in postoperative mortality rate between Asian studies and Western studies for the 90-day mortality risk (4.4% vs 6.8%; P = .02). A lower mortality risk in Asian studies was also found for perihilar cholangiocarcinoma.⁵⁰ The patient characteristics differ among regions, and a higher incidence of obesity is observed in Western populations. This adds to co-morbidity scores and increases the surgical risk. It remains unclear to what extent the higher postoperative mortality in Western studies is explained by differences in baseline patient and tumor characteristics or differences in patient care.

Intrahepatic cholangiocarcinoma patients often present with multifocal disease, for which guidelines recommend palliative chemotherapy rather than surgical resection.^{51–54} Hepatic arterial infusion pump (HAIP) has been investigated for patients with unresectable iCCA confined to the liver. The HAIP delivers high doses of chemotherapy with floxuridine directly in the hepatic artery via a surgically implantable pump. A recent study found that patients with multifocal iCCA had a similar median OS after HAIP floxuridine chemotherapy compared to surgical resection, whereas major complication (6% vs 25%) and postoperative 30-day mortality (1% vs 6%) were lower in the HAIP group.⁵⁵ These results suggest consideration of HAIP instead of surgical resection in patients with multifocal iCCA, in particular in patients with an increased surgical risk.

Several limitations should be addressed for this systematic review and meta-analysis. First, the included studies were all retrospective studies that are prone to selection bias and information bias with underreporting of postoperative complications. A high risk of bias was detected at the level of 'clear reporting of clinical information.' This was mainly because most studies (24/32) lacked details on the ASA classification. However, resection type was specified in most studies (24/32), which was relevant for the subgroup analyses. Postoperative morbidity and mortality are increased in patients with a poor future liver remnant function and volume. The results of the included studies were too heterogeneous for analyzing such risk factors. Furthermore, many studies include patients over a long inclusion period because of the rarity of the disease, of which some included patients from the 1990s. Since 1990, many advancements in technique and surgical management were introduced over time. The year of surgery was not reported in any of the included studies. A subgroup analysis, however, found that mortality was similar in studies with only patients after 2000. However, a difference in postoperative mortality could not be demonstrated. Finally, some of the pooled analyses included a smaller number of studies and patients resulting in wider 95% CIs.

In conclusion, the best estimate for the 90-day postoperative mortality after resection of iCCA is 6.1% with a major complication rate of 22.2%. Ninety-day postoperative mortality rates were lower in Asian studies compared to Western centers (4.4% vs 6.8%). When informing patients or comparing outcomes across hospitals, post-operative mortality rates after liver resection should be reported for 90-days due to the significant number of patients that may survive the first 30 days (ie, PHLF) but will succumb within the following 60 days, with consideration of the diagnosis and extent of liver resection.

Funding/Support

This research did not receive any specific funding from any agencies in the public, commercial, or not-for-profit areas.

Conflict of interest/Disclosure

The authors have no conflicts of interests or disclosures to report.

ARTICLE IN PRESS

A.-M. van Keulen et al. / Surgery xxx (2022) 1–10

Acknowledgments

The authors wish to thank Sabrina Meertens-Gunput from the Erasmus MC Medical Library for developing and updating the search strategies.

References

- DeOliveira ML, Cunningham SC, Cameron JL, et al. Cholangiocarcinoma: thirtyone-year experience with 564 patients at a single institution. *Ann Surg.* 2007;245:755–762.
- Fitzmaurice C, Dicker D, Pain A, et al. Global Burden of Disease Cancer Collaboration. The Global Burden of Cancer 2013. JAMA Oncol. 2015;1:505–527.
- 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.
- Buettner S, van Vugt JL, JN IJ, Groot Koerkamp B. Intrahepatic cholangiocarcinoma: current perspectives. Onco Targets Ther. 2017;10:1131–1142.
- Rizvi S, Khan SA, Hallemeier CL, Kelley RK, Gores GJ. Cholangiocarcinoma evolving concepts and therapeutic strategies. *Nat Rev Clin Oncol.* 2018;15: 95–111.
- 6. Kirstein MM, Vogel A. Epidemiology and risk factors of cholangiocarcinoma. *Visc Med.* 2016;32:395–400.
- **7.** Alvaro D, Bragazzi MC, Benedetti A, et al. Cholangiocarcinoma in Italy: a national survey on clinical characteristics, diagnostic modalities and treatment. Results from the "Cholangiocarcinoma" committee of the Italian Association for the Study of Liver Disease. *Dig Liver Dis.* 2011;43:60–65.
- Squires MH, Cloyd JM, Dillhoff M, Schmidt C, Pawlik TM. Challenges of surgical management of intrahepatic cholangiocarcinoma. *Expert Rev Gastroenterol Hepatol.* 2018;12:671–681.
- **9.** de Jong MC, Nathan H, Sotiropoulos GC, et al. Intrahepatic cholangiocarcinoma: an international multi-institutional analysis of prognostic factors and lymph node assessment. *J Clin Oncol.* 2011;29:3140–3145.
- Farges O, Fuks D, Boleslawski E, et al. Influence of surgical margins on outcome in patients with intrahepatic cholangiocarcinoma: a multicenter study by the AFC-IHCC-2009 study group. Ann Surg. 2011;254:824–829;discussion 30.
- 11. Ribero D, Pinna AD, Guglielmi A, et al. Surgical approach for long-term survival of patients with intrahepatic cholangiocarcinoma: a multi-institutional analysis of 434 patients. *Arch Surg.* 2012;147:1107–1113.
- 12. Mavros MN, Economopoulos KP, Alexiou VG, Pawlik TM. Treatment and prognosis for patients with intrahepatic cholangiocarcinoma: systematic review and meta-analysis. *JAMA Surgery*. 2014;149:565–574.
- **13.** Ma KW, Cheung TT, She WH, et al. Major postoperative complications compromise oncological outcomes of patients with intrahepatic cholangiocarcinoma after curative resection A 13-year cohort in a tertiary center. *Asian J Surg.* 2019;42:164–171.
- Lang H, Sotiropoulos GC, Sgourakis G, et al. Operations for intrahepatic cholangiocarcinoma: single-institution experience of 158 patients. J Am Coll Surg. 2009;208:218–228.
- Zhang XF, Bagante F, Chakedis J, et al. Perioperative and long-term outcome for intrahepatic cholangiocarcinoma: impact of major versus minor hepatectomy. *J Gastrointest Surg.* 2017;21:1841–1850.
- Hobeika C, Cauchy F, Poté N, et al. Short- and long-term outcomes of liver resection for intrahepatic cholangiocarcinoma associated with the metabolic syndrome. World J Surg. 2019;43:2048–2060.
- Rafecas A, Torras J, Fabregat J, et al. Intrahepatic cholangiocarcinoma: prognostic factors for recurrence and survival in a series of 67 patients treated surgically at a single center. *Cir Esp.* 2020;99:506–513.
- Shubert CR, Habermann EB, Truty MJ, Thomsen KM, Kendrick ML, Nagorney DM. Defining perioperative risk after hepatectomy based on diagnosis and extent of resection. J Gastrointest Surg. 2014;18:1917–1928.
- **19.** Liberati A, Altman DG, Tetzlaff J, et al. The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate healthcare interventions: explanation and elaboration. *BMJ*. 2009;339:b2700.
- Moola SMZ, Tufanaru C, Aromataris E, et al. Adelaide: The Joanna Briggs Institute; 2017.
- Conci S, Viganò L, Ercolani G, et al. Outcomes of vascular resection associated with curative intent hepatectomy for intrahepatic cholangiocarcinoma. *Eur J Surg Oncol.* 2020;46:1727–1733.
- **22.** Morimoto Y, Tanaka Y, Ito T, et al. Long-term survival and prognostic factors in the surgical treatment for intrahepatic cholangiocarcinoma. *J Hepatobiliary Pancreat Surg.* 2003;10:432–440.
- 23. Schnitzbauer AA, Eberhard J, Bartsch F, et al. The MEGNA score and preoperative anemia are major prognostic factors after resection in the German intrahepatic cholangiocarcinoma cohort. Ann Surg Oncol. 2020;27:1147–1155.
- 24. Buettner S, Ten Cate DWG, Bagante F, et al. Survival after resection of multiple tumor foci of intrahepatic cholangiocarcinoma. *J Gastrointest Surg.* 2019;23: 2239–2246.
- 25. Spolverato G, Yakoob MY, Kim Y, et al. Impact of complications on long-term survival after resection of intrahepatic cholangiocarcinoma. *Cancer*. 2015;121:2730–2739.

- **26.** Reames BN, Ejaz A, Koerkamp BG, et al. Impact of major vascular resection on outcomes and survival in patients with intrahepatic cholangiocarcinoma: a multi-institutional analysis. *J Surg Oncol.* 2017;116:133–139.
- Merath K, Chen Q, Bagante F, et al. A multi-institutional international analysis of textbook outcomes among patients undergoing curative-intent resection of intrahepatic cholangiocarcinoma. *JAMA Surg.* 2019;154:e190571.
- Olthof PB, Elfrink AKE, Marra E, et al. Volume-outcome relationship of liver surgery: a nationwide analysis. Br J Surg. 2020;107:917-926.
- 29. Lee GC, Gamblin TC, Fong ZV, et al. Facility type is associated with margin status and overall survival of patients with resected intrahepatic cholangiocarcinoma. *Ann Surg Oncol.* 2019;26:4091–4099.
- **30.** Liu H, Cen X, Suo T, et al. Trends and hospital variations in surgical outcomes for cholangiocarcinoma in New York state. *World J Surg.* 2017;41:525–537.
- Jutric Z, Johnston WC, Hoen HM, et al. Impact of lymph node status in patients with intrahepatic cholangiocarcinoma treated by major hepatectomy: a review of the National Cancer Database. *HPB (Oxford)*. 2016;18:79–87.
- Filmann N, Walter D, Schadde E, et al. Mortality after liver surgery in Germany. Br J Surg. 2019;106:1523–1529.
- Bartsch F, Tripke V, Baumgart J, Hoppe-Lotichius M, Heinrich S, Lang H. Extended resection of intrahepatic cholangiocarcinoma: a retrospective singlecenter cohort study. Int J Surg. 2019;67:62–69.
- Beetz O, Weigle CA, Cammann S, et al. Preoperative leukocytosis and the resection severity index are independent risk factors for survival in patients with intrahepatic cholangiocarcinoma. *Langenbecks Arch Surg.* 2020;405:977–988.
- Bektas H, Yeyrek C, Kleine M, et al. Surgical treatment for intrahepatic cholangiocarcinoma in Europe: a single center experience. J Hepatobiliary Pancreat Sci. 2015;22:131–137.
- **36.** Bergeat D, Sulpice L, Rayar M, et al. Extended liver resections for intrahepatic cholangiocarcinoma: friend or foe? *Surgery*. 2015;157:656–665.
- Guglielmi A, Ruzzenente A, Campagnaro T, et al. Intrahepatic cholangiocarcinoma: prognostic factors after surgical resection. World J Surg. 2009;33:1247–1254.
- Lurje G, Bednarsch J, Czigany Z, et al. The prognostic role of lymphovascular invasion and lymph node metastasis in perihilar and intrahepatic cholangiocarcinoma. *Eur J Surg Oncol.* 2019;45:1468–1478.
- Nickkholgh A, Ghamarnejad O, Khajeh E, et al. Outcome after liver resection for primary and recurrent intrahepatic cholangiocarcinoma. *BJS Open.* 2019;3: 793–801.
- Cho SY, Park SJ, Kim SH, et al. Survival analysis of intrahepatic cholangiocarcinoma after resection. Ann Surg Oncol. 2010;17:1823–1830.
- **41.** Li H, Wu JS, Wang XT, et al. Major hepatectomy is a safe modality for the treatment of intrahepatic cholangiocarcinoma in selected patients complicated with cirrhosis. *J Gastrointest Surg.* 2014;18:194–199.
- 42. Luo X, Yuan L, Wang Y, Ge R, Sun Y, Wei G. Survival outcomes and prognostic factors of surgical therapy for all potentially resectable intrahepatic cholangiocarcinoma: a large single-center cohort study. J Gastrointest Surg. 2014;18:562–572.
- Miyata T, Yamashita YI, Yamao T, et al. Prognostic impacts of postoperative complications in patients with intrahepatic cholangiocarcinoma after curative operations. Int J Clin Oncol. 2017;22:526–532.
- **44.** Si A, Li J, Yang Z, et al. Impact of anatomical versus non-anatomical liver resection on short- and long-term outcomes for patients with intrahepatic cholangiocarcinoma. *Ann Surg Oncol.* 2019;26:1841–1850.
- Yoh T, Hatano E, Nishio T, et al. Significant improvement in outcomes of patients with intrahepatic cholangiocarcinoma after surgery. World J Surg. 2016;40:2229–2236.
- 46. Zhu Y, Song J, Xu X, Tan Y, Yang J. Safety and feasibility of laparoscopic liver resection for patients with large or multiple intrahepatic cholangiocarcinomas: a propensity score based case-matched analysis from a single institute. *Medicine (Baltimore)*. 2019;98:e18307.
- An S-L, Liu L-G, Rong W, et al. Surgical outcome and prognostic factors in intrahepatic cholangiocarcinoma: a single-center experience of 114 cases. Int J Clin Exp Med. 2017;10:1156–1163.
- Hammond JS, Guha IN, Beckingham IJ, Lobo DN. Prediction, prevention and management of postresection liver failure. *Br J Surg.* 2011;98:1188–1200.
- Truant S, El Amrani M, Skrzypczyk C, et al. Factors associated with fatal liver failure after extended hepatectomy. *HPB (Oxford)*. 2017;19:682–687.
- Olthof PB, Miyasaka M, Koerkamp BG, et al. A comparison of treatment and outcomes of perihilar cholangiocarcinoma between Eastern and Western centers. *HPB (Oxford)*. 2019;21:345–351.
- Amin MB, Greene FL, Edge SB, et al. The eighth edition AJCC Cancer Staging Manual: continuing to build a bridge from a population-based to a more "personalized" approach to cancer staging. CA Cancer J Clin. 2017;67:93–99.
- Bridgewater J, Galle PR, Khan SA, et al. Guidelines for the diagnosis and management of intrahepatic cholangiocarcinoma. J Hepatol. 2014;60: 1268–1289.
- Weber SM, Ribero D, O'Reilly EM, Kokudo N, Miyazaki M, Pawlik TM. Intrahepatic cholangiocarcinoma: expert consensus statement. *HPB (Oxford)*. 2015;17:669–680.
- Massani M, Nistri C, Ruffolo C, et al. Intrahepatic chemotherapy for unresectable cholangiocarcinoma: review of literature and personal experience. Updates Surg. 2015;67:389–400.
- Franssen S, Soares KC, Jolissaint JS, et al. Comparison of hepatic arterial infusion pump chemotherapy vs resection for patients with multifocal intrahepatic cholangiocarcinoma. JAMA Surg. 2022;157:590–596.