vein embolization and staging laparoscopy should be considered in selected patients. Additional evidence is needed to fully define the role of orthotopic liver transplant. Portal and lymph node involvement worsen the prognosis and long-term survival, and surgery is the only option that can lengthen it. Improvements in adjuvant therapy are essential for improving long-term outcome. Furthermore, the lack of effective chemotherapy drugs and radiotherapy approaches leads us to consider R1 resection as an option, because operated patients have a longer survival rate than those who did not undergo surgery.

**Key words:** Cholangiocarcinoma; Klatskin tumor; Outcome; Pronostic factors; Survival rate

**Core tip:** Klatskin described the specific clinical characteristics in 1965, and the tumor is often referred to as Klatskin tumor. Cholangiocarcinomas (CC) are the second most frequent primary hepatic malignancy. Hilar cholangiocarcinoma (HCC) is the most common type, and accounts for approximately 60% to 67% of all cholangiocarcinoma cases. There is not a staging system that permits us to compare all series and extract some conclusions to increase the long-survival rate in this dismal disease. Neither the extension of resection, according to the sort of HCC, is a closed topic. Some authors defend limited resection (mesohepatectomy with S1, S1 plus S4b-S5, local excision for papillary tumours, etc.) while others insist in the compulsory removal of an extended hepatic resection with portal vein bifurcation removal to reach cure. As there is not an ideal adjuvant therapy, R1 resection can be justified to prolong the survival rate. Morbidity and mortality rates changed along the last decade, but variability is the rule, with morbidity and mortality rates ranging from 14% to 76% and from 0% to 19%, respectively. Conclusion: Surgical resection continues to be the main treatment of HCC. Negative resection margins achieved with major hepatic resections are associated with improved outcome. Pre-resectional management with biliary drainage, portal vein embolization and staging laparoscopy should be considered in selected patients. Additional evidence is needed to fully define the role of orthotopic liver transplant. Portal and lymph node involvement worsen the prognosis and long-term survival, and surgery is the only option that can lengthen it. Improvements in adjuvant therapy are essential for improving long-term outcome. Furthermore, the lack of effective chemotherapy drugs and radiotherapy approaches leads us to consider R1 resection as an option, because operated patients have a longer survival rate than those who did not undergo surgery.

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**Abstract**

Cholangiocarcinomas are the second most frequent primary hepatic malignancy, and make up from 5% to 30% of malignant hepatic tumors. Hilar cholangiocarcinoma (HCC) is the most common type, and accounts for approximately 60% to 67% of all cholangiocarcinoma cases. There is not a staging system that permits us to compare all series and extract some conclusions to increase the long-survival rate in this dismal disease. Neither the extension of resection, according to the sort of HCC, is a closed topic. Some authors defend limited resection (mesohepatectomy with S1, S1 plus S4b-S5, local excision for papillary tumours, etc.) while others insist in the compulsory removal of an extended hepatic resection with portal vein bifurcation removed to reach cure. As there is not an ideal adjuvant therapy, R1 resection can be justified to prolong the survival rate. Morbidity and mortality rates changed along the last decade, but variability is the rule, with morbidity and mortality rates ranging from 14% to 76% and from 0% to 19%, respectively. Conclusion: Surgical resection continues to be the main treatment of HCC. Negative resection margins achieved with major hepatic resections are associated with improved outcome. Pre-resectional management with biliary drainage, portal vein embolization and staging laparoscopy should be considered in selected patients. Additional evidence is needed to fully define the role of orthotopic liver transplant. Portal and lymph node involvement worsen the prognosis and long-term survival, and surgery is the only option that can lengthen it. Improvements in adjuvant therapy are essential for improving long-term outcome. Furthermore, the lack of effective chemotherapy drugs and radiotherapy approaches leads us to consider R1 resection as an option, because operated patients have a longer survival rate than those who did not undergo surgery.

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**Core tip:** Klatskin described the specific clinical characteristics in 1965, and the tumor is often referred to as Klatskin tumor. Cholangiocarcinomas (CC) are the second most frequent primary hepatic malignancy. Hilar cholangiocarcinoma (HCC) is the most common type, and accounts for most of CC cases. These tumors are slowly growing, and have a tendency to local spread and infrequent distant metastases. The most common presentation is with jaundice. The majority of HCC are small infiltrating tumors. Long-term survival in patients with HCC depends critically on complete tumor resection. This work is an important update concerning outcome of surgical management in Klatskin tumors.

**Introduction**

Carcinomas arising from the confluence of the hepatic ducts were first described by Altemeier et al[8]. Klatskin[2]
described the specific clinical characteristics in 1965, and the tumor is often referred to as Klatskin tumor. Cholangiocarcinomas (CC) are the second most frequent primary hepatic malignancy and make up from 5% to 30% of malignant hepatic tumors. Hilar cholangiocarcinoma (HCC) is the most common type, and accounts for approximately 50% to 60% of all CC cases (intrahepatic, hilar and distal)\[11-13\]. These tumors are slowly growing, and have a tendency to local spread and infrequent distant metastases. The most common presentation is with the onset of jaundice. The majority of HCC are small infiltrating tumors. Approximately 90% of malignant-appearing hilar strictures prove to be HCC\[8\].

Adenocarcinoma is the most common histologic subtype. Three morphologic subtypes of cholangiocarcinoma have been described: sclerosing (70%), nodular (20%), and papillary (5%)\[6\]. Characteristics of nodular and sclerosing types may coexist.

Long-term survival in patients with HCC depends critically on complete tumor resection. In the absence of widespread disease, the likelihood of achieving a complete resection requires examination of all factors related to local tumor extent, which increasingly has become critically on complete tumor resection. In the absence of widespread disease, the likelihood of achieving a complete resection requires examination of all factors related to local tumor extent, which increasingly has become

| Tumor, nodes and metastases definitions |
|----------------------------------------|
| Primary tumor                          |
| Tis Carcinoma in situ                  |
| T1 Tumor confined to the bile duct histologically |
| T2 Tumor invades beyond the wall of the bile duct |
| T3 Tumor invades the liver, gallbladder, pancreas, and/or ipsilateral branches of the portal vein or hepatic artery |
| T4 Tumor invades any of the following: main portal vein or its branches bilaterally, common hepatic artery, or other adjacent structures, such as the colon, stomach, duodenum, or abdominal wall. |
| Regional lymph nodes                   |
| N0 No regional lymph node metastasis   |
| N1 Regional lymph node metastases      |
| Metastasis                             |
| M0 No distant metastasis               |
| M1 Distant metastasis                  |

| Stage grouping                         |
|----------------------------------------|
| Stage I Tis, N0, M0                     |
| Stage IA T1, N0, M0                     |
| Stage IB T2, N0, M0                     |
| Stage IIA T3, N0, M0                    |
| Stage IIB T1, N1, M0                    |
| Stage III T4, any N, M0                 |
| Stage IV Any T, any N, M1               |

| Stage | Hilar involvement | Portal vein | Lobar atrophy |
|-------|-------------------|-------------|---------------|
| T1    | Biliary confluence ± 1/2 unilateral extension to second-order biliary radicles | No | No |
| T2    | Biliary confluence ± unilateral extension to second-order biliary radicles | Ipsilateral | Ipsilateral |
| T3    | Biliary confluence ± unilateral extension to second-order biliary radicles | Yes/No | Yes/No |
| T4    | Biliary confluence ± unilateral extension to second-order biliary radicles | Yes/No | Yes/No |
| T5    | Biliary confluence ± unilateral extension to second-order biliary radicles | Yes/No | + Contrateral |
| T6    | Biliary confluence ± unilateral extension to second-order biliary radicles | Yes/No | + Contrateral |
| T7    | Biliary confluence ± unilateral extension to second-order biliary radicles with contralateral hepatic lobar atrophy; | Bilateral | Yes/No |
| T8    | Biliary confluence ± unilateral/ bilaterally | | |

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| T5    | Biliary confluence ± unilateral extension to second-order biliary radicles | Yes/No | Yes/No |
| T6    | Biliary confluence ± unilateral extension to second-order biliary radicles | Yes/No | + Contrateral |
| T7    | Biliary confluence ± unilateral extension to second-order biliary radicles with contralateral hepatic lobar atrophy; | Bilateral | Yes/No |
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The TNM staging system of the American Joint Commission on Cancer (AJCC) (Table 1) is the most commonly used for staging of HCC. However, this system is based on histological criteria and does not provide information on the potential for resectability. de Jong et al\[9\] conclude that the AJCC T-classification criteria did not stratify patients with regard to prognosis and that depth of tumor invasion is a better predictor of long-term outcome. Besides that, the histologic type of tumor may also modify the staging and type of surgery required\[10\].

Therefore, other staging systems have been used to predict resectability and evaluate the extent of resection. The modified Bismuth-Corlette (B-C) classification stratifies patients according to the extent of biliary involvement by tumor\[11-13\]. Although it does not incorporate radial tumor extension, it provides a useful preoperative terminology to describe the extent of the hepatic resection that will be necessary to encompass the longitudinal intraductal extension of HCC.

The preoperative clinical T-stageing system of the Memorial Sloan Kettering Cancer Centre (MSKCC) (Table 2), as proposed by Jarnagin and Blumgart (MSKCC), defines both the longitudinal and radial extension of HCC, which are critical factors in the determination of resectability\[14,15\]. This staging system incorporates three factors based on preoperative imaging studies: (1) location and extent of ductal involvement; (2) presence or absence of portal vein invasion; and (3) presence or absence of hepatic lobar atrophy. Criteria for unresectable disease include: locally advanced tumor extending to secondary biliary radicles bilaterally, unilateral segmental bile ducts with contralateral portal vein branch involvement, encasement or occlusion of the main portal vein proximal to its bifurcation, and atrophy of one hepatic lobe with contralateral tumor extension to segmental bile ducts. Of note, the right bile duct is shorter and therefore more likely to be involved when the tumor appears at the confluence. Patients who have distant metastases, including metastases to lymph node groups beyond the hepatoduodenal ligament are considered unresectable. The modified Bismuth-Corlette (B-C) classification system defines the extent of biliary involvement by tumor\[11-13\] and is based on preoperative imaging studies: (1) location and extent of ductal involvement; (2) presence or absence of portal vein invasion; and (3) presence or absence of hepatic lobar atrophy. Criteria for unresectable disease include: locally advanced tumor extending to secondary biliary radicles bilaterally, unilateral segmental bile ducts with contralateral portal vein branch involvement, encasement or occlusion of the main portal vein proximal to its bifurcation, and atrophy of one hepatic lobe with contralateral tumor extension to segmental bile ducts. Of note, the right bile duct is shorter and therefore more likely to be involved when the tumor appears at the confluence. Patients who have distant metastases, including metastases to lymph node groups beyond the hepatoduodenal ligament are considered unresectable.

**STAGING AND RESECTABILITY**

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Characteristics of the growth pattern of HCC include: transmural invasion of bile ducts, radial extension into periductal tissue and adjacent structures, and longitudinal extension along the bile ducts in the submucosa\cite{18}. The papillary phenotype is associated with better prognosis\cite{3}. In contrast, longitudinal spread along the duct will with microscopic submucosal extension is characteristic of mass-forming and periductal-infiltrating subtypes; this biologic feature often impedes obtaining histologically negative margins\cite{19}. These tumors are often accompanied by both direct and lymphatic invasion into the periductal tissues, causing marked fibrosis and infiltration of inflammatory cells. These histologic changes give a macroscopic similarity between the tumor and peritumoral inflammatory changes that make preoperative and intraoperative biopsies diagnostically challenging. Radial extension of HCC is also common, often resulting in invasion of the portal vein, hepatic arteries and the hepatic parenchyma adjacent to the hilar plate.

When analysing survival according to staging, Li et al\cite{19} in their audit of 215 patients found that the results from univariate analyses suggest that histological grade, lymph node metastasis, vascular invasion, neuroinvasion, R1 resection and T2 or T3 stage were significant predictors for poor survival rates; by multivariate analysis, only lymph node metastasis and R1 resection were significantly associated with poor survival rates.

Series with more than 100 cases in consulted literature are scarce, and those ones that fulfil this condition cover a prolonged period of time and are retrospective. The resectability rates were highly variable, ranging between 28% and 95%, and curative resection rates ranged between 14% and 95%\cite{4,14,15,17,20-42}. Such wide variability of resectability is probably due to heterogeneous methods of patient selection, differences in preoperative imaging techniques, and the broad range of data for inclusion in these studies. The report of DeOliveira et al\cite{43} where 282 HCC patients are assessed, is one of the biggest published series of only one institution, together with that one of Nagoya group, but it covers a 31-year period and is retrospective\cite{44}. Apart from the changes in management over the course of a long period of time, as can be seen in Figure 1, the resectability rate in that study was 62% and R0 resection was achieved only in 19% of cases\cite{43}.

Even in high-volume centres, the resectability rate is about 30% of all patients with HCC, with the operative mortality rate ranging from 0% to 15%. After curative resection, the 1-, 3- and 5-year survival rates range from 50% to 70%, 30% to 40%, and 10% to 40%, respectively (Figure 1)\cite{14,20,43-48}.

The major determinants of resectability include ex-

\begin{figure}
\centering
\includegraphics[width=\textwidth]{figure1.png}
\caption{Changes in pre-, intra-, and postoperative management over the course of the study period (With permission. Courtesy of Professor Nimura). ENBD: Endoscopic naso-biliary drainage; MDCT: Multidetector-row computed tomography; PTBD: Percutaneous transhepatic biliary drainage; PTCS: Percutaneous transhepatic cholangioscopy.}
\end{figure}
tent of vascular invasion, hepatic lobar atrophy, amount of hepatic parenchyma involved, and extent of spread within the biliary tree. Hepatic lobar atrophy with contralateral portal vein or hepatic artery encasement or contralateral tumor extension to secondary biliary radicles may preclude resection. Bilateral hepatic disease and presumed insufficient hepatic reserve preclude resection. Even with current imaging technology, accurate determination of tumor resectability pre-operatively may occur in as few as 60%-74% of patients\(^{46,54}\). Thus, a number of patients undergoing resection with curative intent will be left with a resultant R1 margin status.

**ADJUVANT TREATMENT**

Adjuvant therapy for CC has not been supported by clinical evidence. Recently, gemcitabine has been shown to be active, with response rates of 8%-60% and median survival of 6-16 mo. Therefore, further studies of gemcitabine and of 5-FU plus cisplatin are warranted. For HCC, Cheng et al\(^{[51]}\) reported better survival for patients with Bismuth types III/IV tumors who received adjuvant radiotherapy after curative resection. Todoroki et al\(^{[51]}\) also showed a statistically significance of radiotherapy for R1 radical resection of stage IVa HCC. Thus, radiotherapy is potentially beneficial in patients with positive resection margins or unresectable tumors. However, Vern-Gross concluded that there is no benefit with adjuvant therapy in postoperative setting\(^{[52]}\).

**PREOPERATIVE BILIARY DRAINAGE**

The role of preoperative biliary drainage (PBD) in jaundiced patients remains controversial\(^{22,45,53,54}\). Actually, most patients undergo biliary drainage prior to referral for resection, despite the lack of data showing a benefit. Clearly, the presence of cholangitis mandates biliary decompression, but there is no proof that routine biliary drainage in all patients facilitates resection or reduces postsurgical morbidity\(^{[53,56]}\). On the contrary, the available data would suggest that biliary stents are associated with greater postoperative infection complications\(^{[57,58]}\). Previous studies investigating this issue have been criticized for several design flaws, and whether major hepatic resection in the face of biliary obstruction is associated with a greater risk of liver failure or other complications remains an open question\(^{[59]}\).

Cherqui et al\(^{[60]}\) reported the results of major hepatobiliary resection without PBD in 20 patients with biliary cancer. Postoperative liver failure rate was 5%, and mortality was documented in the same patients.

PBD is associated with an increased risk of cholangitis and prolonged postoperative hospital stay, and can impede the ability to determine the extent of tumor during surgery. Cholangitis after PBD has been reported in 20%-60% of cases and may compromise subsequent surgery with patient dropout. Intraoperative bile cultures have been found to be positive in 65% of patients with PBD, while the rate was 8% in patients without PBD. This may be associated with increased postoperative infections such as wound or intraperitoneal abscesses\(^{[60]}\).

However, unrelieved biliary obstruction is associated with hepatic and renal dysfunction and coagulopathy. Most patients with HCC will benefit from PBD of remnant liver to increase post-resection hypertrophy ability. Reported complications in transhepatic percutaneous catheter placement include: haemobilia, pseudoaneurysm of hepatic artery, fistula between hepatic artery and bile duct or between hepatic artery and portal vein, and catheter tract implantation metastases.

Some randomized controlled trials have revealed that biliary diversion does not improve perioperative results and increases infectious complications. But, also, none of these trials has managed to clarify the safety of major hepatic resection for cholestatic patients with HCC\(^{[53,54]}\). The report of Laurent et al\(^{[61]}\) states some conditions to avoid PBD: onset of jaundice < 2-3 wk, total bilirubin < 200 µmol/L, functional remnant liver (FRL) > 40%, neither endoscopic retrograde cholangiopancreatography nor percutaneous transhepatic cholangiography, and no sepsis. Although the results are not modified for not to drain, in agreement with other authors, undrained patients have a higher postoperative morbidity rate and transfusion requirements, and both facts are important factors of tumor recurrence. Thus, it may depend on each group’s experience to determine whether to use PBD or not. It will be taken into consideration that, if the conditions described by Laurent et al\(^{[61]}\) are not fulfilled, there will be more perioperative transfusion and morbidity if the patient is not drained, which could affect the overall survival and disease-free survival rate.

**PORTAL VEIN EMBOLIZATION**

Resection greater than 80% of total liver volume is associated with major complications and prolonged hospital stay in patients with normal liver function, and resection greater than 60% is associated with an increase of major complications, postoperative hepatic insufficiency and mortality in patients with impaired liver function due to chronic liver disease, chronic biliary obstruction or high-dose chemotherapy\(^{[61,65]}\). Preoperative portal vein embolization (PVE) was first described in 1986 and is currently used to increase FRL volume and function\(^{[65]}\).

Randomized controlled trials and individual institutional series support the safety and efficiency of preoperative PVE\(^{[20,61,65-69]}\). A potential disadvantage of PVE is that it may be difficult to determine preoperatively whether a right or left hepatectomy will be required if the tumor is located centrally in the hilum. At present, there is no evidence to support the routine use of PVE for HCC, but PVE should be considered for potentially resectable patients with normal liver function when anticipated FRL is less than 20% of the total liver volume, or for patients with compromised liver function when
anticipated FRL is less than 40% of the total liver volume. Most patients with HCC present with jaundice and are considered to have cholestasis-induced compromised liver function. There are not many data on the impact and real volume of PVE on FRL liver function increase, associated or not to biliary drainage. Only in the second period in which their series is divided do Cannon et al. use PVE in 9.1% of cases, which means 4.5% out of a total of 110 patients, and despite that use they achieve only 62% of R0 resections.

As a consequence, PVE must be assessed and chosen with caution to avoid the frightening postoperative hepatic insufficiency, one of the main causes of mortality in these patients. Also, its application must be evaluated in accordance with a previous surgical plan, which, if uncertain, could lead us to use another type of tactic, such as associating liver partition and portal vein ligation for staged hepatectomy (ALPPS)\(^\text{[5]}\).

**LIVER RESECTION: HOW MUCH IS ENOUGH?**

In the last 20 years the use of hepatic resection in patients with HCC has risen. The objective of all the techniques and of the tendency to major resection with or without resection of vessels is to obtain free resection margins. The 5-year survival rate in patients undergoing non-curative resection for HCC is below 10%\(^\text{[1]}\). The 5-year survival rate for operated patients is with curative intention 11%-41% (Figure 1). All scientific community agrees that surgical resection is the only potentially curative treatment for CC, but the disease is usually advanced at the time of diagnosis and mostly treated by chemoradiotherapy or palliative therapy, including biliary drainage or stenting. Resectability rates are low because of early infiltration of the tumor into adjacent structures such as hepatic artery, portal vein and caudate lobe. In patients treated with curative intent, an extended hemihepatectomy is often needed to achieve negative margins. Preoperative jaundice and extended procedures are important risk factors for postoperative complications\(^\text{[3]}\).

The aims of surgery in HCC are: (1) to achieve macroscopic removal of the tumor; (2) to restore satisfactorily the bile flow to the gut; and (3) to minimize postoperative liver failure or death. At the beginning of last decade, resection was possible only in 20% of cases, and the operative mortality was 10%. The median survival was only 20 mo, but resected patients enjoyed a good quality of life\(^\text{[4]}\). Last decade saw an aggressive approach to HCC with an increasing use of major hepatic resections\(^\text{[1,4,10-12,15,17-19]}\). The resectability rate increased to 80% with the addition of hepatic resection to bile duct resection without increasing the postoperative death rate. Bismuth et al.\(^\text{[2]}\) and Pichlmayr et al.\(^\text{[3]}\) suggested a stagewise management strategy with the prime objective of achieving complete surgical resection of the tumor without leaving behind macroscopic residual disease. Patients with Bismuth types I and II were treated by bile duct resection. For Bismuth stage IIa/IIb lesions, resection of the corresponding hemiliver was recommended. However, major hepatic resection is a formidable operation in patients with a cholestatic liver and carries a high complication rate, with a morbidity of up to 81% and mortality rates of between 6% and 10% in the most advanced centres.

Vascular encasement with or without biliary obstruction may result in segment or lobar atrophy. Long-standing biliary obstruction can cause moderate atrophy, whereas concomitant portal venous compromise usually produces rapid and severe atrophy of the involved segments\(^\text{[9]}\). Approximately 30% of patients subjected to surgical exploration show evidence of lobar atrophy\(^\text{[10]}\). It is one of the problems of the PVE, together with vascular involvement not detected before embolization.

The caudate lobe is frequently involved by either direct invasion or ductal extension. Caudate bile ducts can drain to both the right and left hepatic ducts; in fact, some series have identified microscopic tumor infiltration into the caudate lobe in nearly all patients with HCC\(^\text{[21]}\). In general, the primary drainage of the caudate lobe is into the left hepatic duct\(^\text{[78]}\). For this reason, it has been alleged the necessity to resect the caudate lobe in Bismuth type II from now on.

Ikeyama et al.\(^\text{[10]}\), in their audit of 54 consecutive type I and II HCC resected patients, concluded that for nodular and infiltrating tumors right hepatectomy is essential; for papillary tumors, bile duct resection with or without limited hepatectomy is adequate. But the problem is that it is very difficult to know these issues preoperatively and intraoperatively. Nuzzo et al.\(^\text{[3]}\) reached the same conclusion in their audit of 440 patients, showing that pathologic factors independently predicted overall and disease-free survival at multivariate analysis.

Major hepatic resections have increased the proportion of R0 resections\(^\text{[6,14,39,77]}\), improved the outcome of disease-free survival, and decreased the prevalence of hepatic recurrence\(^\text{[79]}\). Surgical results improved in the 1990s thanks to a better ability to perform R0 resections, which is likely due to increasing use of major hepatic resection and portal resections, as well as the improvement of preoperative management concerning both prognosis and FRL preparation and care\(^\text{[74,78]}\). Recent studies have also reported an improvement in morbidity and mortality in comparison with previous decades, which probably responds to advances in overall perioperative care. Also, the improvement of preoperative management has had a consequence, as can be seen in the report of Nagino et al.\(^\text{[44]}\).

Nonetheless, it is uncertain whether the major hepatic resection may improve the survival of patients with B-C types I or II HCC. Ikeyama et al.\(^\text{[10]}\) and Jang et al.\(^\text{[10]}\) showed survival benefit in right hepatectomy with caudate lobectomy for nodular and sclerosing tumors, but not for papillary ones. However, others have reported a non-significant difference between hepatectomy and isolated bile duct resection in B-C types I and II tumors\(^\text{[77]}\).
Regarding proximal margin, it can be stated nowadays that survival outcomes improve when bile duct resection is associated with hepatectomy, even in patients with B-C types I and II tumors\cite{14,26}. In the series published by Jar-nagin et al.\cite{14,26} in 2001, the 5-year survival was 37% when a hepatic resection was performed (84% of R0 resections) and 0% when only a bile duct excision was performed (56% of R0 resections). The best results are obtained with a right hepatectomy, probably because this surgical technique facilitates en-bloc resection of the tumor and surrounding tissues and thereby increases radicality\cite{26}. In the series of Neuhaus et al.\cite{24,25}, the worst outcomes after hepatectomy with curative intent were obtained in patients undergoing left hepatectomy. Although Nimura defended the radical surgery of left-sided Klatskin tumors by performing a left trisectionectomy, this is characterized by high morbidity rates and by mortality rates superior to 10\%\cite{78,79}. The analysis of recurrence after R0 resection with hepatectomy shows a low frequency of local recurrence, but a high frequency of peritoneal seeding recurrence\cite{26}. Then, manipulation of the tumor as well as biopsies may favor local recurrence, and this is the reason why some authors advise en-bloc resection including surrounding vessels, a “non-touch technique”, in order to avoid this cause of recurrence.

The hepatectomy must include the caudate lobe, since this is a frequent site of tumor recurrence when it is not included in the resection piece. However, as it happens with other “evidences” related to Klatskin tumor treatment, there are no controlled studies that support this recommendation\cite{32,80}. Performing a perioperative biopsy of the biliary resection margin in the liver remnant is common practice for most surgeons.

In a recent report of Ribero et al.\cite{81}, in the analysis of 82 cases, the group of patients who had primary R0 was compared with those patients who achieved a secondary R0 after an intraoperative additional resection, and also with the patients who were R1. The 1-, 3- and 5-year survival rates were similar in the groups with primary R0 and secondary R0, but different in R1 patients (5-year survival rate: 50\%, 30.8\% and 0\% respectively). The authors concluded that an additional resection of a positive proximal bile duct margin, albeit associated with an increased risk of biliary fistula, offers a significant survival benefit and should be attempted whenever possible. But this Italian group does not re-operate on those patients who the pathologist changes to R1 resection in the postoperative study, and thus, although they only have 13 cases that underwent re-resection, they do not defend re-operations on patients when this occurs. However, it is necessary to take into consideration that frozen biopsy is often not concluding and that resection extension, when the biopsy is positive, is frequently impracticable\cite{24}. This explains why perioperative biopsies in this location have low profitability. Furthermore, such resection of margin-positive proximal duct does improve survival even when a negative margin can be achieved with additional resection\cite{82}.

**LYMPHATIC SPREAD**

In addition to extension along the bile ducts, HCC often metastasizes via the lymphatics. Lymphatic metastases are found in 30\% to 50\% of patients undergoing resection\cite{14,43,34}. Hilar and pericholedochal lymph nodes (LN) are the most commonly involved, followed by periportal, common hepatic, posterior pancreaticoduodenal, celiac and preaortic ones\cite{83}. Metastasis in regional LN is an important prognostic factor that affects survival after the resection of an HCC\cite{30}. Kitagawa et al.\cite{24} evaluated 110 patients that underwent resection for HCC with LN dissection, including both the regional and para-aortic ones, and found that 47\% of patients had no involved LN, 35\% had metastases in regional LN and 17\% had metastases in regional and para-aortic LN. The 5-year survival was 36\% for patients with negative LN, 15\% for patients with metastases in regional LN and 12\% for patients with metastases in regional and para-aortic LN. Other studies have reported a worse survival in patients with LN involvement beyond the hepatoduodenal ligament, with a 5-year survival rate ranging from 0\% to 6\%\cite{45}. Consequently, routine LN dissection beyond hepatoduodenal ligament is not recommended. Patients with macroscopically involved LN beyond hepatoduodenal ligament are considered to have unresectable disease, even though some surgeons resect them if they find them intraoperatively.

Only one study has presented the number of affected LN as a variable than worsens survival\cite{86}.

**VASCULAR RESECTION**

Radial growth of the tumor may infiltrate the surrounding vessels. Right hepatic artery involvement is more frequent due to its proximity to the biliary bifurcation. Contralateral artery infiltration to the hepatic resection that is to be performed is a reason for contraindication of surgical treatment. Portal involvement is present in 20\%-30\% of R0 resections and its preoperative identification is achieved with a precision of 85\%. In the experience of Nagoya University, in approximately one third of the patients whose portal vein is resected because of apparent infiltration, this is not histologically confirmed\cite{46}. However, most of these patients had a tumor infiltration adjacent to the vein, and the margin would have been positive without vein resection. On the other hand, vascular resection was not associated with a significant increase of morbimortality. Anyhow, resection can improve survival in some patients when R0 resection is achieved.

Encasement or occlusion of the main portal vein or vessels supplying the hepatic remnant is considered a contraindication to surgery\cite{14}. Recent reports have shown that en-bloc resection with vascular reconstruction can achieve negative margins with a 10\% perioperative mortality in selected patients.

Portal vein resection and reconstruction has been carried out in HCC with conflicting results\cite{24,87}. Although several retrospective series have not shown difference in
operative mortality between the patients that underwent portal vein resection and those that did not [25], the impact of portal vein resection on long-term survival is less clear. Neuhaus et al [30] proposed portal vein resection as part of a “non-touch” resection of the tumor and surrounding tissue. Portal vein resection was identified as a positive independent prognostic factor in their multivariate analysis of patients undergoing R0 resections, when mortality within the first 60 d was excluded. Nevertheless, overall mortality within 60 d after portal vein resection was 17%, in comparison with 5% in patients without portal resection, and all the deaths occurred after non-curative resections. Other studies have reported similar or worse survival in patients undergoing portal vein en-bloc resection [23-25]. The role of routine portal vein resection (as stated by Neuhaus) is not likely to be clearly designed unless a randomized clinical trial is completed. However, Hemmings rejects the routine performance of this procedure and in 2012 the Nagoya group reported a 5-year survival rate of 40% in the last period of portal resection, but a morbidity of 57.3% [26].

Portal resection must be recommended whenever the tumor cannot be freed from it, since the microscopic invasion of the portal vein does not seem to influence on survival when a vascular resection is carried out, whereas the macroscopic invasion does have negative results on survival.

Nishio et al [9] concluded that although lymph node metastasis and macroscopic portal vein involvement were independent negative prognostic factors, the 5-year survival rate obtained in patients with portal vein resection or lymph node metastasis still was about 10% (Table 3). Even in patients with both cancer invasion of the portal vein and regional lymph node metastasis, or with para-aortic lymph node metastasis, curative resection resulted in significantly longer survival than the one found in unresected patients.

Some groups had 100% morbidity and mortality in arterial resections, although in arterial and portal combined resections mortality was 43%, and the overall percentage of positive margins was 32% [9]. de Jong et al [9] reported in a recent paper that combined hepatectomy, extrahepatic biliary duct resection and portal vein resection can offer long-term survival in some patients with advanced HCC, with 17.6% mortality rate and 28% 5-year survival rate.

Some authors recommend hepatectomy with simultaneous arterial and portal vein resection. They reach 66% of R0 resection with 2% mortality rate, 54% morbidity rate and 1-, 3-, and 5-year survival rates of 78.9%, 36.3%, and 30.3%, respectively, but these data are not reproducible [9].

Su et al [3], Miyazaki et al [30] and Muñoz et al [31] reported as a conclusion that, although both portal vein and hepatic artery resection are independent poor prognostic factors after curative operative resection for locally advanced HCC, portal vein resection is acceptable from an operative risk perspective and might improve the prognosis in the selected patients, but combined hepatic artery resection cannot be justified because the 3-year survival rate is 9%.

## LIVER TRANSPLANT

Orthotopic liver transplant (OLT) is contraindicated in HCC because of disappointing long-term outcomes. However, a recent multi-institutional study in the United States, including 280 patients with earlier-stage tumors who received aggressive neoadjuvant chemoradiation, has reported that transplantation remarkably improves survival: the 1- and 5-year survival rates were 74% and 38%, respectively [9]. The Mayo Clinic protocol sets a strict selection of the patients candidates to liver transplant. Although the selection is highly rigorous and biased for patients with biologically favourable disease, the early results published by the Mayo group showed an 82% 5-year survival rate [9]. The histological analysis of resected pieces confirmed N0 and R0 state in all the patients. However, only 58% of the patients had histologically confirmed cancer.

Liver transplantation is currently done only in the setting of clinical trials. It offers the advantage of resection of all structures that may be involved by the tumor, including portal vein, bilateral hepatic ducts and atrophic hepatic lobes. Thus, total hepatectomy may permit R0 resection even in locally advanced tumors, which are beyond resection criteria. Efficacy of neoadjuvant therapy and transplantation is demonstrated by comparing results with the natural history of the disease. Untreated HCC

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**Table 3 Prognostic factors and 5-year survival rate**

| Hilary cholangiocarcinoma | No. of patients | Prognostic factors | Operative mortality(%) | 5-yr SV (%) |
|---------------------------|-----------------|--------------------|------------------------|------------|
| Jarnagin et al [23]        | 80              | Margin, hepatectomy, differentiation | 10 | 27 |
| Seyama et al [79]          | 58              | Lymph nodes        | 0 | 40 |
| Dinant et al [75]          | 99              | Margin, resection period, lymph nodes | 15 | 27 |
| DeOliveira et al [31]      | 281             | Margin, lymph nodes | 5 | 10 |
| Rea et al [47]             | 46              | Lymph nodes, tumor grade, bilirubin | 9 | 26 |
| Silva et al [55]           | 45              | Tumor stage, margin | 9 | 11 |
| Witzigmann et al [10]      | 60              | Residual tumor status, grading | 8 | 22 |
| Batton et al [48]          | 59              | Chemotherapy, margin, lymph nodes | 5 | 20 |
| Wahab et al [90]           | 243             | Margin, S1 resection, lymph nodes, grading | 7 | 16 |
| de Jong et al [30]         | 305             | Lymph nodes, margin | 5 | 20 |

SV: Survival rate.
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Table 4 Morbidity and mortality rate and R0 resections

| Ref.       | Resections | RO (%) | Morbidity | Mortality  | 5-yr Survival |
|------------|------------|--------|-----------|------------|---------------|
| Burke et al| 30     | 83     | NA        | 6          | 45            |
| Nakach et al| 109   | 26     | 47        | 4          | 11            |
| Jarnagin et al| 80   | 78     | 64        | 10         | 26            |
| Nimmura et al| 55    | 84     | 41        | 6          | 41            |
| Jarnagin et al| 106  | 77     | 62        | 8          | NA            |
| Seyama et al| 87    | 64     | 43        | 0          | 40            |
| Kossuue et al| 65    | 52     | 37        | 9          | 33            |
| Neuhaus et al| 80   | 61     | 55        | 8          | 22            |
| Launois et al| 131  | NA     | NA        | 19         | NA            |
| Kondo et al| 40     | 95     | 48        | 0          | NA            |
| Dinant et al| 99    | 31     | 66        | 15         | 27            |
| Gerhards et al| 112  | 14     | 65        | 18         | 19            |
| Hemming et al| 53   | 80     | 40        | 9          | 35            |
| Ijitsma et al| 42   | 65     | 76        | 12         | 19            |
| Kawarada et al| 65   | 64     | 28        | 2.3        | 26            |
| Klemmaxer et al| 151  | 77     | NA        | 10         | 28            |
| Miyazaki et al| 76   | 71     | 33        | 13         | 26            |
| Nimmura et al| 142  | 61     | 49        | 9          | 26            |
| Su et al[44] | 49    | 49     | 47        | 10         | 15            |
| Todoroki et al| 101  | 14     | 14        | 4          | 28            |
| DeOliveira et al| 281  | 62     | 60        | 5          | 30            |
| Nagino et al| 574   | 76.5   | 43.1      | 4.7        | 32.5          |
| Ren et al[8] | 46    | 80     | 52        | 9          | 26            |
| Nuazzo et al| 440   | 77.3   | 47.5      | 8.6        | 25.5          |
| Ito et al[4] | 38    | 63     | 32        | 0          | 33            |
| Kawasaki et al| 79   | 68     | 14        | 1.3        | 22            |

NA: Not available.

has a 50%-70% mortality rate within 12 mo, which is much worse than 55% 5-year survival for patients who entered the Mayo Clinic protocol and 71% 5-year survival after transplantation[9,10].

The Cincinnati Transplant Tumor Registry reported 28% 5-year survival, with a tumor recurrence rate of 51%[28]. The Spanish liver transplant centres provided similar results, with 30% 5-year survival rate and 53% tumor recurrence rate, in 36 patients with unresectable, non-disseminated HCC[9]. As a consequence of such initial results and the limited availability of organs, HCC was perceived as a relative contraindication to OLT. Also, it is a well-known fact that 55% of HCC even in T2 stages have affected LN, which is one of the contraindications to transplant[3].

A further complication to transplants in HCC is that, as response to postoperative radiotherapy and chemotherapy is low both in R1 and recurrence, tumors must have a more favourable biological behaviour, and if sizes bigger than 2 cm are rejected for rescue with liver transplant, then very few patients can be candidates to be transplanted[3]. It is important to remember that, out of the 281 cases analysed by DeOliveira, 58% were > 2 cm and hilar involvement occurred in 28%[3].

Schüle et al[9] concluded that an acceptable survival rate could be achieved by transplantation for HCC with LN metastases as the only exclusion criterion, even if they use living donors. In this article, the authors got a 5-year survival rate of 50% in those patients with negative LN.

Nowadays, OLT cannot be considered as a standard therapy for HCC in patients with resectable disease, but it offers a potential option to patients with underlying primary sclerosing cholangitis. Additional studies are necessary to define the role of OLT in depth.

MORBIDITY AND MORTALITY

Due to the complex biliary and liver resections required to obtain complete tumor removal, the risks of perioperative morbidity and mortality are significant. Morbidity and mortality rates range from 14% to 76% and from 0% to 19%, respectively. Perioperative morbidity includes haemorrhage, biliary fistula, hepatic insufficiency and infectious complications. Among them, infectious complications are particularly common and account for 50% to 80% of all complications[11,42]. The postoperative liver failure and its morbidity have been joined with the extension of hepatic resection[11]. However, recent publications suggest a decrease in morbidity and mortality with the use of preoperative PVE, even in extended hepatectomies[13,42,61,59] (Table 4).

OUTCOME OF RESECTION

Published 5-year survival rates range from 25% to 40% in recent series, and, even, it has been reported that many clinical and histological factors have a positive impact on long-term outcome, including negative histologic margin status[99,100], concomitant hepatic resection[8], absence of nodal involvement[14,23,48,101], low TNM status[102], well-differentiated tumor grade[42], papillary tumor morphology[46,44,84], and lack of perineural invasion[39]. Complete resection with negative histologic margins is the only modifiable factor and, for that reason, the primary aim of surgical therapy. There is a close association between hepatic resection and negative margins[42]. The effect of R1 resection vs no resection on outcome has been object of discussion and analysis in surgical literature, with some recent studies that report improvement in survival after R1 resection in comparison with patients with resectable disease[42].

Recurrence after resection occurs quite frequently, in up to 50%-75% of cases[10,22,74]. The median recurrence time ranges from 12 to 43 mo[10,22,47,6]. Prognostic factors for recurrence-free survival include histologic grade, T and N stages, and margin status[10,22,76,102]. Since patients with recurrent disease are not candidates for curative therapy, advances in adjuvant therapy are essential to improve long-term outcome. However, the effectiveness of radiotherapy and chemotherapy is still very limited. In the report of Cherqui et al[9], the authors concluded that adjuvant radiotherapy was not associated with an improvement in long-term overall survival in patients with resected HCC.

CONCLUSION

Surgical resection continues to be the main treatment of HCC. Negative resection margins enhanced by major
hepatic resections are associated with improved outcome. Pre-resectional management with biliary drainage, PVE and staging laparoscopy should be considered in selected patients. Additional evidence is needed to fully define the role of OLT. Improvements in adjuvant therapy are essential for improving long-term outcome. Portal and node involvement worsens the prognosis and long-term survival, and surgery is the only option that can lengthen it. Furthermore, the lack of effective chemotherapy and radiotherapy treatments, at this moment, leads us to consider R1 resection as an option because these patients have a longer survival rate than patients who do not undergo resection.

REFERENCES

1 Altermeyer WA, GALL EA, ZINNINGER MM, HOXWORTH PJ. Sclerosing carcinoma of the major intrahepatic bile ducts. Ama Arch Surg 1957; 75: 450-460: discussion 460-461 [PMID: 13457619 DOI: 10.1001/archsurg.1957.00800101015]

2 Klatskin G. Adenocarcinoma of the Hepatic Duct at Its Bifurcation Within the Porta Hepatis. An Unusual Tumor with Distinctive Clinical and Pathological Features. Am J Med 1965; 38: 241-256 [PMID: 14256720 DOI: 10.1016/0002-9343(65)90178-6]

3 Burke EC, Jarnagin WR, Hochwald SN, Pisters PW, Fong Y, Blumgart LH. Hilar Cholangiocarcinoma: patterns of spread, the importance of hepatic resection for curative operation, and a presurgical clinical staging system. Ann Surg 1998; 228: 385-394 [PMID: 9742921 DOI: 10.1097/00000658-199809000-00011]

4 Nakeeb A, Pitt HA, Sohn TA, Coleman J, Abrams RA, Piantadosi S, Hruban RH, Lillemoe KD, Yeo CJ, Cameron JL. Cholangiocarcinoma. A spectrum of intrahepatic, perihilar, and distal tumors. Ann Surg 1996; 224: 463-473; discussion 473-476 [PMID: 8857851 DOI: 10.1097/00000658-199610000-00005]

5 Buc E, Lesurtel M, Belghiti J. Is preoperative histological diagnosis necessary before referral to major surgery for cholangiocarcinoma? HPB (Oxford) 2008; 10: 98-105 [PMID: 18773064 DOI: 10.1016/j.hpb.2007.08.015]

6 Weinbren K, Mutum SS. Pathological aspects of cholangiocarcinoma. J Pathol 1983; 139: 217-238 [PMID: 6293844 DOI: 10.1002/path.1711390210]

7 Hanno LE, Greatrex KV, Bach AM, Fong Y, Blumgart LH. Cholangiocarcinoma at the hepatic hilus: sonographic findings. AJR Am J Roentgenol 1997; 168: 985-989 [PMID: 9124155 DOI: 10.2214/ajr.168.4.9124155]

8 Schwartz LH, Coakley FV, Sun Y, Blumgart LH, Fong Y, Panicek DM. Neoplastic pancreaticobiliary duct obstruction: evaluation with breath-hold MR cholangiopancreatography. AJR Am J Roentgenol 1998; 170: 1491-1495 [PMID: 9696160 DOI: 10.2214/ajr.170.6.9696160]

9 de Jong MC, Hong SM, Augustine MM, Goggins MG, Wolffgang C, Hirose K, Schullik RD, Choti MA, Anders RA, Pawlik TM. Hilar cholangiocarcinoma: tumor depth as a predictor of outcome. Arch Surg 2011; 146: 697-703 [PMID: 21690466 DOI: 10.1001/archsurg.2011.122]

10 Ikeyama T, Nagino M, Oda K, Ebata T, Nishio H, Nimura Y. Surgical approach to bismuth Type I and II hilar cholangiocarcinomas: audit of 54 consecutive cases. Ann Surg 2007; 246: 1052-1057 [PMID: 18043110 DOI: 10.1097/SLA.0b013e3181a42f97]

11 Bismuth H. Corlette MB. Intrahepatic cholangioanastomotic anastomosis in carcinoma of the hilus of the liver. Surg Gynecol Obstet 1975; 140: 170-178 [PMID: 1079096]

12 Bismuth H, Castaing D, Traynor O. Resection or palliation: priority of surgery in the treatment of hilar cancer. World J Surg 1988; 12: 39-47 [PMID: 2449769 DOI: 10.1007/BF01658484]

13 Bismuth H, Nakache R, Diamond T. Management strategies in resection for hilar cholangiocarcinoma. Ann Surg 1992; 215: 31-38 [PMID: 1309988 DOI: 10.1097/00000658-199201000-00005]

14 Jarnagin WR, Fong Y, DeMatteo RP, Gonen M, Burke EC, Bodniewicz BS J, Youssef BA M, Klimstra D, Blumgart LH. Staging, resectability, and outcome in 225 patients with hilar cholangiocarcinoma. Ann Surg 2001; 234: 507-517; discussion 517-519 [PMID: 11573044 DOI: 10.1097/00000658-200106000-00010]

15 Nimura Y, Hayakawa N, Kaniya J, Kondo S, Shionoya S. Hepatic segmentectomy with caudate lobe resection for bile duct carcinoma of the hepatic hilus. World J Surg 1990; 14: 535-543; discussion 544 [PMID: 2166381 DOI: 10.1007/BF01658686]

16 Hayashi S, Miyazaki M, Kondo Y, Nakajima N. Invasive growth patterns of hepatic hilar ductal carcinoma. A histologic analysis of 18 surgical cases. Cancer 1994; 73: 2922-2929 [PMID: 8198989]

17 Jarnagin WR, Bowne W, Klimstra DS, Ben-Porat L, Roggin K, Cymes K, Fong Y, DeMatteo RP, D’Angelica M, Koea J, Blumgart LH. Papillary phenotype confers improved survival after resection of hilar cholangiocarcinoma. Ann Surg 2005; 241: 703-712; discussion 712-714 [PMID: 15849506 DOI: 10.1097/01.sla.0000168017.94472.fd]

18 Shimada H, Niimoto S, Matsuha A, Nakagawa G, Kobayashi M, Tsuchiya S. The infiltration of bile duct carcinoma along the bile duct wall. Int Surg 1988; 73: 87-90 [PMID: 2840410]

19 Li H, Qin Y, Cui Y, Chen H, Hao X, Li Q. Analysis of the surgical outcome and prognostic factors for hilar cholangiocarcinoma: a Chinese experience. Dig Surg 2011; 28: 226-231 [PMID: 21540611 DOI: 10.1159/000327361]

20 Seyama Y, Kubota K, Sano K, Noite T, Takayama T, Kusoge T, Makuchii M. Long-term outcome of extended hemihepatectomy for hilar bile duct cancer with no mortality and high survival rate. Ann Surg 2003; 238: 73-83 [PMID: 12863968 DOI: 10.1097/01.sla.0000074960.55004.72]

21 Mizumoto R, Kawarada Y, Suzuki H. Surgical treatment of hilar carcinoma of the bile duct. Surg Gynecol Obstet 1986; 162: 153-158 [PMID: 3945893]

22 Lai EC, Mok FP, Fan ST, Lo CM, Chu KM, Liu CL, Wong J. Preoperative endoscopic drainage for malignant obstructive jaundice. Br J Surg 1994; 81: 1195-1198 [PMID: 7741850 DOI: 10.1002/bjs.1808010859]

23 Kosuge T, Yamamoto J, Shimada K, Yamasaki S, Makuchii M. Improved surgical results for hilar cholangiocarcinoma with no postoperative mortality and no positive ductal margins: results of a prospective study. Ann Surg 2005; 241: 703-712; discussion 712-714 [PMID: 15849506 DOI: 10.1097/01.sla.0000168017.94472.fd]

24 Bismuth H, Corlette MB. Intrahepatic cholangioanastomotic anastomosis in carcinoma of the hilus of the liver. Surg Gynecol Obstet 1975; 140: 170-178 [PMID: 1079096]

25 Landin S, Gerhards MF, Rauws EA, Busch OR, Gouma DJ,
van Gulik TM. Improved outcome of resection of hilar cholangiocarcinoma (Klatskin tumor). Ann Surg Oncol 2006; 13: 872-880 [PMID: 16614876 DOI: 10.1245/ASCO.2006.05.053]

Gazzaniga GM, Filardo M, Bagarolo C, Mori L. Surgery for hilar cholangiocarcinoma: an Italian experience. J Hepatobiliary Pancreat Surg 2000; 7: 122-127 [PMID: 10982603 DOI: 10.1007/s005340051605]

Gerhards MF, van Gulik TM, de Wit LT, Obertop H, Gouma DJ. Evaluation of morbidity and mortality after resection for hilar cholangiocarcinoma—a single center experience. Surgery 2000; 127: 395-404 [PMID: 10776430 DOI: 10.1067/msy.2000.104250]

Hadjis NS, Blennkhorn JI, Alexander N, Benjamin IS, Blumgart LH. Outcome of radical surgery in hilar cholangiocarcinoma. Surgery 1990; 107: 597-604 [PMID: 2162862]

Hasegawa S, Ikai I, Fujii H, Hatano E, Shimahara Y. Surgical resection of hilar cholangiocarcinoma: analysis of survival and postoperative complications. World J Surg 2007; 31: 1256-1263 [PMID: 17455285 DOI: 10.1007/s00268-007-0001-y]

Hemming AW, Reed AI, Fujita S, Foley DP, Howard RJ. Surgical management of hilar cholangiocarcinoma. Ann Surg 2005; 241: 693-699; discussion 699-702 [PMID: 18349505 DOI: 10.1016/s0003-4975(01)08945-2]

Ijitsma AJ, Appelants BM, de Jong KP, Porte RJ, Peeters PM, Slooff MJ. Extrahepatic bile duct resection in combination with liver resection for hilar cholangiocarcinoma: a report of 42 cases. J Gastrointest Surg 2004; 8: 686-694 [PMID: 15585329 DOI: 10.1016/j.gassur.2004.04.006]

Kawarada Y, Das BC, Nagamura T, Tabata M, Taoka H. Surgical treatment of bile duct hilar carcinoma: experience with 25 consecutive hepatectomies. J Gastrointest Surg 2002; 6: 617-624 [PMID: 12127130 DOI: 10.1016/s1091-255x(01)00088-7]

Klemprnauer J, Rirdder GJ, von Wasielewski R, Werner M, Weimann A, Pichlmayr R. Resectional surgery of hilar cholangiocarcinoma: a multivariate analysis of prognostic factors. J Clin Oncol 1997; 15: 947-954 [PMID: 9060532]

Lee SG, Lee YJ, Park KM, Hwang S, Min PC. One hundred and eleven liver resections for bile duct cancer. J Hepato-Pancreat Surg 2000; 7: 135-141 [PMID: 10982605 DOI: 10.1007/s005340051617]

Miyazaki M, Ito H, Nakagawa K, Ambiru S, Shimizu H, Shimizu Y, Kato A, Nakamura S, Momoto H, Nakajima N, Kimura F, Suwa T. Aggressive surgical approaches to hilar cholangiocarcinoma: hepatic or local resection? Surgery 1998; 133: 131-136 [PMID: 9814139 DOI: 10.1016/s0039-6060(98)00249-1]

Nimura Y, Kamiya J, Kondo S, Nagino M, Uesaka K, Oda K, Sano T, Yamamoto H, Hayakawa N. Aggressive preoperative management and extended surgery for hilar cholangiocarcinoma: Nagoya experience. J Hepato-Pancreat Surg 2000; 7: 155-162 [PMID: 10982608 DOI: 10.1007/s005340050170]

Su CH, Tsay SH, Wu CC, Shyr YM, King KL, Lee CH, Lui WY, Liu TJ, Peng FK. Factors influencing postoperative morbidity, mortality, and survival after resection for hilar cholangiocarcinoma. Arch Surg 1996; 131: 136-141 [PMID: 8363917 DOI: 10.1001/archsurg.131.2.136]

Takada T, Amano H, Yasuda H, Nimura Y, Matsushiro T, Kato H, Nagakawa T, Nakayama T. Is postoperative adjuvant chemotherapy useful for gallbladder carcinoma? A phase III multicenter prospective randomized controlled trial in patients with resected pancreaticobiliary carcinoma. Cancer 2002; 95: 1685-1695 [PMID: 12365016 DOI: 10.1002/cncr.110831]

Todori T, Kamamoto T, Koi K, Takehashi H, Yoshida S, Kashiwagi H, Takada Y, Otsuka M, Fukao K. Radical resection of hilar bile duct carcinoma and predictors of survival. Br J Surg 2000; 87: 306-313 [PMID: 10718799 DOI: 10.1046/j.1365-2168.2000.01343.x]
Serrablo A et al. Outcome of surgical resection in Klatskin tumors.

MG. Adenocarcinoma of the distal bile duct. A clinicopathologic outcome analysis after curative resection. *Dig Surg* 2000; 17: 36-41 [PMID: 10728380 DOI: 10.1159/00018798]

57 Nimura Y. Preoperative biliary drainage before resection for cholangiocarcinoma (Prev). *HPB* (Oxford) 2008; 10: 133-145 [PMID: 18735400 DOI: 10.1016/j.ipal.2009.211626]

59 Laurent A, Tayar C, Cherqui D. Cholangiocarcinoma: preoperative biliary drainage (Con). *HPB* (Oxford) 2008; 10: 126-129 [PMID: 18770920 DOI: 10.1016/j.ipal.2009.210474]

94 Lieser MJ, Barry MK, Rowland C, Istrup DM, Nagorney DM. Surgical management of intrahepatic cholangiocarcinoma: a 31-year experience. *J Hepatobiliary Pancreat Surg* 1999; 5: 41-47 [PMID: 9883753 DOI: 10.1007/FL00099449]

96 Seyama Y, Mukuchu M. Current surgical treatment for bile duct cancer. *World J Gastroenterol* 2007; 13: 1505-1515 [PMID: 17461441]

100 Hemming AW, Reed AJ, Howard RJ, Fujita S, Hochwald SN, Cardi JG, Hawkins IF, Vauthey JN. Preoperative portal vein embolization for extended hepatectomy. *Ann Surg* 2003; 237: 668-691; discussion 691-693 [PMID: 12724655 DOI: 10.1097/01.sla.0000065256.16728:CO]

103 Shoup M, Gonen M, D’Angelaica M, Jarnagin WR, DeMatteo RP, Schwartz LH, Tuorto S, Blumgart LH, Fong Y. Volumetric analysis predicts hepatic dysfunction in patients undergoing major liver resection. *J Gastrointest Surg* 2003; 7: 325-330 [PMID: 12654556 DOI: 10.1016/S1091-255X(02)00370-4]

106 Vauthey JN, Chaux A, Do KA, Bilimoria MM, Fenstermacher MJ, Charmsangavaj C, Hicks M, Asfassler G, Lauwers G, Hawkins IF, Cardi J. Standardized measurement of the future liver remnant prior to extended liver resection: methodology and clinical associations. *Surgery* 2000; 127: 512-519 [PMID: 10819059 DOI: 10.1067/msy.2000.105294]

108 Shirabe K, Shimada M, Gion T, Hasegawa H, Takenaka K, Utsunomiya T, Sugimachi K. Postoperative liver failure after major hepatic resection for hepatocellular carcinoma in the modern era with special reference to remnant liver volume. *J Am Coll Surg* 1999; 188: 304-309 [PMID: 10058520 DOI: 10.1016/S1072-7515(98)00391-9]

110 Ebata T, Nagino M, Kamiya J, Uesaka K, Nagasaka T, Nimura Y. Hepatocyte with portal vein resection for hilar cholangiocarcinoma: audit of 52 consecutive cases. *Ann Surg* 2003; 238: 720-727 [PMID: 14578735 DOI: 10.1097/01.sla.0000048447.16651.7B]

112 Nagino M, Kamiya J, Nishio H, Ebata T, Arai T, Nimura Y. Two hundred forty consecutive portal vein embolizations before extended hepatectomy for biliary cancer: surgical outcome and long-term follow-up. *Ann Surg* 2006; 243: 364-372 [PMID: 16495702 DOI: 10.1016/s0002-1482.11867.14]

113 Farges O, Belghiti J, Kianmanesh R, Regimbeau JM, Santoro R, Vilgrain V, Denys A, Sauvanet A. Portal vein embolization for extended hepatectomy. *Ann Surg* 2001; 233: 720-727 [PMID: 12650779 DOI: 10.1097/00000542-199612000-00018]

119 Sano T, Shimada K, Sakamoto Y, Yamamoto J, Yamasaki S, Kosuge T. One hundred two consecutive hepatic resections for perihilar cholangiocarcinoma with zero mortality. *Ann Surg* 2006; 244: 240-247 [PMID: 16858166 DOI: 10.1097/01.sla.000019389.65619.38]

127 Vauthey JN, Pawlik TM, Abdalla EK, Arens JF, Nemr RA, Verbeek PC, van Leeuwen DJ, de Wit LT, Reeders JW, Smits NJ, Bosma A, Huijbregts K, van der Heyde MN. Benign ﬁbrosis of the proximal bile duct in hilar cholangiocarcinoma. *Ann Surg* 2003; 237: 208-217 [PMID: 12560779 DOI: 10.1097/00000542-199611000-00018]

130 Shingu Y, Ebata T, Arai T, Kamiya J, Santoro R, D’Amico DF, Langer AM, Belli G, Bresadola F, Calise F, Dalla Valle R, Fichet D, Ferruzzi G, Giordano A, Gori A, Gouarderes J, Grommes R, Hasegawa Y, Hermanns E, Igami T, Ishii M, Jarnigan W, Pitt H, Gores G, Busuttil R, Pappas T, Nagino M, Kamiya J, Uesaka K, Sano T, Yama moto H, Hayakawa N, Nimura Y. Lymph node metastasis from hilar cholangiocarcinoma: audit of 110 patients who underwent regional and paraaortic node dissection. *Ann Surg* 2001; 233: 385-392 [PMID: 11224627 DOI: 10.1097/00000542-199612000-00013]

140 Nuzzo G, Giuliani F, Ardiffo F, Giovannini I, Aldrighetti L, Berrisi S, Cavallari M, Charnsangavej C, Hicks M, Alsfasser G, Lauwers G, Hawkins IF, Cardi J. Standardized measurement of the future liver remnant prior to extended liver resection: methodology and clinical associations. *Surgery* 2000; 127: 512-519 [PMID: 10819059 DOI: 10.1067/msy.2000.105294]

147 Ebata T, Nagino M, Kamiya J, Uesaka K, Nagasaka T, Nimura Y. Hepatocyte with portal vein resection for hilar cholangiocarcinoma: audit of 52 consecutive cases. *Ann Surg* 2003; 238: 720-727 [PMID: 14578735 DOI: 10.1097/01.sla.0000048447.16651.7B]

149 Shingu Y, Ebata T, Nishio H, Igami T, Shimoyama H, Nimura Y. Lymph node metastasis from hilar cholangiocarcinoma: audit of 110 patients who underwent regional and paraaortic node dissection. *Ann Surg* 2001; 233: 385-392 [PMID: 11224627 DOI: 10.1097/00000542-199612000-00013]

156 Shingu Y, Ebata T, Arai T, Kamiya J, Santoro R, D’Amico DF, Langer AM, Belli G, Bresadola F, Calise F, Dalla Valle R, Fichet D, Ferruzzi G, Giordano A, Gori A, Gouarderes J, Grommes R, Hasegawa Y, Hermanns E, Igami T, Ishii M, Jarnigan W, Pitt H, Gores G, Busuttil R, Pappas T, Nagino M, Kamiya J, Uesaka K, Sano T, Yamamoto H, Hayakawa N, Nimura Y. Lymph node metastasis from hilar cholangiocarcinoma: audit of 110 patients who underwent regional and paraaortic node dissection. *Ann Surg* 2001; 233: 385-392 [PMID: 11224627 DOI: 10.1097/00000542-199612000-00013]

161 Nuzzo G, Giuliani F, Ardiffo F, Giovannini I, Aldrighetti L, Berrisi S, Cavallari M, Charnsangavej C, Hicks M, Alsfasser G, Lauwers G, Hawkins IF, Cardi J. Standardized measurement of the future liver remnant prior to extended liver resection: methodology and clinical associations. *Surgery* 2000; 127: 512-519 [PMID: 10819059 DOI: 10.1067/msy.2000.105294]

162 Ebata T, Nagino M, Kamiya J, Uesaka K, Nagasaka T, Nimura Y. Hepatocyte with portal vein resection for hilar cholangiocarcinoma: audit of 52 consecutive cases. *Ann Surg* 2003; 238: 720-727 [PMID: 14578735 DOI: 10.1097/01.sla.0000048447.16651.7B]
Serrablo A et al. Outcome of surgical resection in Klatskin tumors

Ann Surg 2008; 248: 273-279 [PMID: 18650638 DOI: 10.1097/SLA.0b013e318172b8b4]

86 Nagino M, Nimura Y, Nishio H, Ebata T, Igami T, Matsu-shita M, Nishikimi N, Kamei Y. Hepatectomy with simultaneous resection of the portal vein and hepatic artery for advanced perihilar cholangiocarcinoma: an audit of 50 consecutive cases. Ann Surg 2010; 252: 115-123 [PMID: 20531001 DOI: 10.1097/SLA.0b013e3181e463a7]

87 Miyazaki M, Kato A, Itó H, Kimura F, Shimizu H, Ohtsuka M, Yoshidome H, Yoshitomi H, Furukawa K, Nozawa S. Combined vascular resection in operative resection for hilar cholangiocarcinoma: does it work or not? Surgery 2007; 141: 581-588 [PMID: 17462457 DOI: 10.1016/j.surg.2006.09.016]

88 Meyer CG, Penn I, James L. Liver transplantation for cholangiocarcinoma: results in 207 patients. Transplantation 2000; 69: 1633-1637 [PMID: 10836374 DOI: 10.1097/00007890-200004270-00019]

89 Nishio H, Nagino M, Nimura Y. Surgical management of hilar cholangiocarcinoma: the Nagoya experience. HPB (Oxford) 2005; 7: 259-262 [PMID: 18352023 DOI: 10.1080/13651820500205000]

90 de Jong MC, Marques H, Clary BM, Bauer TW, Marsh JW, Ribero D, Majno P, Hatzaras I, Walters DM, Barbas AS, Mega R, Schulick RD, Choti MA, Geller DA, Barroso E, Menta G, Capussotti L, Pawlik TM. The impact of portal vein resection on outcomes for hilar cholangiocarcinoma: a multi-institutional analysis of 305 cases. Cancer 2012; 118: 4737-4747 [PMID: 22415526 DOI: 10.1002/cncr.27492]

91 Muñoz L, Roayaie S, Mamán D, Fishbein T, Shiefer P, Emre S, Miller C, Schwartz ME. Hilar cholangiocarcinoma involving the portal vein bifurcation: long-term results after resection. J Hepatobiliary Pancreat Surg 2002; 9: 237-241 [PMID: 12140613 DOI: 10.1017/s00024610002025]

92 Becker NS, Rodíguez JA, Barshes NR, O’Mahony CA, Goss JA, Aloia TA. Outcomes analysis for 280 patients with cholangiocarcinoma treated with liver transplantation over an 18-year period. J Gastrointest Surg 2008; 12: 117-122 [PMID: 17963015 DOI: 10.1017/s11605-007-0335-4]

93 Heimbach JK, Gores GJ, Haddock MG, Alberts SR, Pedersen R, Kremers W, Nyberg SL, Ishihata MB, Rosen CB. Predictors of disease recurrence following neoadjuvant chemoradiotherapy and liver transplantation for unresectable perihilar cholangiocarcinoma. Transplantation 2006; 82: 1703-1707 [PMID: 17198263 DOI: 10.1097/01.tp.0000253551.43583.d1]

94 Rea DJ, Heimbach JK, Rosen CB, Haddock MG, Alberts SR, Kremers WK, Gores GJ, Nagorney DM. Liver transplantation with neoadjuvant chemoradiation is more effective than resection for hilar cholangiocarcinoma. Ann Surg 2005; 242: 451-458; discussion 458-461 [PMID: 16135931]

95 Rosen CB, Heimbach JK, Gores GJ. Surgery for cholangiocarcinoma: the role of liver transplantation. HPB (Oxford) 2008; 10: 186-189 [PMID: 18773052 DOI: 10.1080/13651820801992542]

96 Robles R, Figuera J, Turrión VS, Margarit C, Moya A, Varo E, Calleja J, Valdivieso A, Valdecasas JC, López P, Gómez M, de Vicente E, Loinaz C, Santoyo J, Fleitas M, Bernardos A, Lladó L, Ramírez P, Bueno FS, Jauretxe E, Parrilla P. Spanish experience in liver transplantation for hilar and peripheral cholangiocarcinoma. Ann Surg 2004; 239: 265-271 [PMID: 14745336 DOI: 10.1097/01.sla.0000108702.45715.81]

97 Kurokawa N, Tsukada K, Hatakeyama K, Muto T. The mode of lymphatic spread in carcinoma of the bile duct. Ann Surg 1996; 172: 239-243 [PMID: 8862075 DOI: 10.1016/s0002-9610(96)00156-0]

98 Schüle S, Altendorf-Hofmann A, Uteß F, Rauchfuß F, Freesmeyer M, Knösel T, Dittmar Y, Settmacher U. Liver transplantation for hilar cholangiocarcinoma—a single-centre experience. Langenbecks Arch Surg 2013; 398: 71-77 [PMID: 23053456 DOI: 10.1017/s0002432233001078]

99 Itó F, Cho CS, Rikkers LF, Weber SM. Hilar cholangiocarcinoma: current management. Ann Surg 2009; 250: 210-218 [PMID: 19638920 DOI: 10.1097/SLA.0b013e3181afe0ab]

100 Kloek J, Ten Kate FJ, Busch OR, Gouma DJ, van Gulik TM. Surgery for extrahepatic cholangiocarcinoma: predictors of survival. HPB (Oxford) 2008; 10: 190-195 [PMID: 18773053 DOI: 10.1080/13651820801992575]

101 Park SW, Park YS, Chung JB, Kang JK, Kim KS, Choi JS, Lee WJ, Kim BR, Song SY. Patterns and relevant factors of tumor recurrence for extrahepatic bile duct carcinoma after radical resection. Hepatogastroenterology 2004; 51: 1612-1618 [PMID: 15532789]

102 Jarnagin WR, Shoup M. Surgical management of cholangiocarcinoma. Semin Liver Dis 2004; 24: 189-199 [PMID: 15192791 DOI: 10.1055/s-2004-828895]

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