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**Maternal choledochal cysts<sup>8</sup> in pregnancy: A systematic review of case reports and case series**

Augustin G *et al.* Maternal choledochal cysts

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**Abstract**

**BACKGROUND**

<sup>9</sup>Choledochal cysts (CC) are cystic dilatations of the biliary tract, usually diagnosed during childhood, with an estimated incidence in the general population of 1:100000. Complications related to CC include rupture, biliary obstruction, and cholangitis. Maternal CC in pregnancy is rarely reported, and there are no guidelines on optimal management.

**AIM**

To systematically review maternal CCs diagnosed during pregnancy or postpartum. We assessed the clinical presentation of CCs, the mode of treatment and delivery, and maternal outcomes.

**METHODS**

A literature search of cases and case series of maternal CC in pregnancy and postpartum was conducted using <sup>13</sup>Medline/PubMed, Web of Science, Google Scholar, and Embase. There were no restrictions on language or publication year. Databases were lastly accessed on September 1, 2022.

## RESULTS

Overall, 71 publications met the inclusion criteria, reporting 97 cases. Eighty-eight were diagnosed during pregnancy and 9 in the puerperium. The most common symptoms were abdominal pain (81.2%) and jaundice (60.4%). Interventions for CC complications were required in 52.5% of the cases, and 34% of pregnancies were induced. Urgent cesarean section (CS) was done in 24.7%. The maternal mortality was 7.2%, while fetal mortality was inconsistently reported, with 8.2%. Cholangitis, CC > 15 cm, and bilirubin levels (> 80 mmol/L) were associated with a higher likelihood of urgent CS and surgical intervention for CC. Bilirubin levels positively correlated with CC size. There was no correlation between age and cyst dimension, gestational age at cyst discovery, and CC size.

## CONCLUSION

Although rare, maternal CC in pregnancy should be included in the evaluation of jaundice with upper abdominal pain. Symptomatology and clinical course are variable, and treatment may range from an expectative approach to emergent surgical CC treatment and urgent CS. While most cases were managed by conservative measures or drainage procedures, CC > 15 cm and progressive cholangitis carry the risk of CC rupture and septic complications, which may increase the rates of unfavorable maternal and fetal outcomes. Therefore, such cases require specific surgical and obstetric interventions.

**Key Words:** Choledochal cyst; Pregnancy; Cholangitis; Surgery; Delivery; Cesarean section

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**Core Tip:** Although rare, maternal choledochal cysts (CC) in pregnancy should be included in the evaluation of jaundice with upper abdominal pain. Symptomatology and clinical course are variable, and treatment may range from an expectative approach to emergent surgical treatment of CC and urgent cesarean section. While most cases were managed by conservative measures or drainage procedures, CC > 15 cm and progressive cholangitis carry the risk of CC rupture and septic complications, which may increase the rates of unfavorable maternal and fetal outcomes. Therefore, such cases require specific surgical and obstetric interventions.

## INTRODUCTION

Choledochal cysts (CC) are cystic dilatations of the biliary tract, usually diagnosed during childhood (80%)<sup>[1,2]</sup>. Todani's classification is based on anatomical characteristics, and it is the most widely used classification method, which divides CCs into five types<sup>[3]</sup>. Type I (cystic dilatation of the common bile duct) is the most common (90%)<sup>[4,5]</sup>. The incidence significantly varies between the East and the West<sup>[1,4]</sup>. The estimated incidence in the general population is 1:100000-1:150000 in Western countries, while in the East, it is much higher, with up to 1:1000 and a male/female ratio of 1:3-4<sup>[1,6]</sup>. Although cholecystitis or choledocholithiasis is more commonly associated with upper right abdominal pain or jaundice, CC should be considered in the differential diagnosis because the treatment approach is different. CC may present with a triad of abdominal pain, jaundice, and a palpable right upper quadrant abdominal mass. CC presentation depends not only on size but on morphology, the presence of sludge/stones, and the status of the remaining biliary tract. When cholangitis develops, the Charcot triad may be present (jaundice, fever with rigors, and right upper quadrant abdominal pain). CC may result in severe complications such as biliary obstruction, cholangitis, or cyst rupture. Furthermore, CC is considered a premalignant condition, another reason for surgical treatment, even in asymptomatic patients<sup>[4,7]</sup>. Gomes *et al*<sup>[8]</sup> made the first report on CC in 1723.

Many theories on the pathogenesis of CC were offered. Some proposed that congenital dysplasia arises from early embryonic development, where dysplasia of the primordial common bile duct (CBD) leads to CBD dilation<sup>[9]</sup>. Another theory is based on a congenital weakness of the bile duct wall that allows cysts to form<sup>[10,11]</sup>. Proponents of the CBD obstruction theory believe that CC forms due to congenital or acquired obstruction of the ducts<sup>[12]</sup>. A so-called dual theory states that cyst formation is due to the weakness of the bile duct wall and obstruction of the ducts<sup>[13]</sup>. However, the most widely accepted theory, the common channel theory, states that the anomalous pancreaticobiliary junction leads to the formation of an abnormally long common channel, which permits reflux of the activated proteolytic pancreatic enzymes, biliary tract wall weakness and, eventually, cystic dilatation<sup>[14]</sup>.

CC treatment in the general population depends on symptomatology and CC features. Most cases require cyst excision, cholecystectomy, and Roux-en-Y hepaticojejunostomy (HJ)<sup>[6,14]</sup>. Urgent indications are cyst rupture, intraluminal biliary bleeding, or severe cholangitis<sup>[12,15]</sup>. Todani's classification determines surgical treatment. Type I and IV should be completely excised, in addition to cholecystectomy and bilioenteric continuity formation. Type II is usually treated with diverticulectomy followed by CBD closure at the diverticulum neck. Endoscopic sphincterotomy is usually sufficient for type III cysts<sup>[11,12,15]</sup>. The treatment of type IV and V is more complex, as both comprise intra- and extrahepatic components, and the intrahepatic component is difficult to treat. When the cyst adheres densely to the portal vein due to an inflammatory reaction, standard full-thickness cyst excision may not be possible. In such cases, the Lilly technique should be considered, which includes curettage or cauterization of cyst mucosa while the serosa remains attached to the portal vein. This resolves cholestasis, and at the same time, it removes the risk of malignant transformation.

Maternal CC in pregnancy is rarely reported, and no studies suggest a possible facilitating impact of pregnancy on CC development or growth. However, the pregnant uterus possibly causes biliary outflow obstruction promoting cystic dilatation, eventually leading to symptoms in prepregnancy asymptomatic individuals<sup>[16]</sup>. In addition, physiological changes during pregnancy can mask different clinical scenarios, including CCs.

The effects of CC and its complications considering pregnancy outcomes have not been studied. Jaundice in pregnancy has potentially severe maternal and fetal consequences, found with intrahepatic cholestasis of pregnancy (ICP) or hemolysis, elevated liver enzymes, low platelets syndrome<sup>[14]</sup>. Adverse fetal outcomes include preterm delivery, respiratory distress, or fetal distress. Therefore, CC with cholestasis and jaundice poses a similar risk to the mother and the fetus as ICP or other cholestatic disorders<sup>[16-18]</sup>.

Due to CC rarity, reliable data on the risk of fetal loss and optimal CC treatment are lacking, and randomized trials or large prospective series do not exist. Therefore, we

aimed to systematically review maternal CCs diagnosed during pregnancy and postpartum and to propose optimal treatment strategies.

## **MATERIALS AND METHODS**

### *Literature search and study selection*

A literature search of cases and case series of maternal CC in pregnancy and postpartum was conducted. We searched MEDLINE/PubMed, Web of Science, Google Scholar, and Embase using a combination of MeSH terms: "maternal", "choledochal cyst", "choledochoceles", "pregnancy", "biliary tract disease", "puerperium" and "postpartum." Puerperium was evaluated up to the 6<sup>th</sup> postpartum week.

The search was limited to human studies, without language or country restrictions, and published until September 2022. Non-English articles were translated and analyzed with the help of medical scientists proficient in the languages mentioned above. We evaluated articles and references from these articles to identify additional cases. Cases with maternal CC treated and diagnosed before pregnancy were excluded. All potentially relevant articles were reviewed by three investigators (AA, GA, and IR), and all disagreements were settled by discussion.

### *Data management*

Data included demographic data (maternal age, parity, gestational week), clinical presentation (pain, jaundice, palpable mass, fever), laboratory data (total bilirubin and liver enzymes), radiological examinations [abdominal ultrasound, abdominal magnetic resonance imaging (MRI), cyst dimensions, cyst type according to Todani's classification], mode of treatment and delivery, maternal and fetal outcome. The risk of bias or study quality was not assessed, as all articles were case reports or small case series. Bilirubin ( $\mu\text{mol/L}$ ) was the only laboratory parameter sufficiently reported to be included in the analysis. If mentioned, the highest diameter of the CC and bilirubin value was used in the calculations. Pregnancies were analyzed through gestational months and trimesters;



the first trimester counts from week zero to the end of the 13<sup>th</sup> week, the second from the 14<sup>th</sup> to the 27<sup>th</sup> week, and the third from the 28<sup>th</sup> to term.

### *Statistical analysis*

Results of descriptive analyses are reported as numbers and proportions in percentages. Inferential statistics were executed on the different numbers of patients depending on the availability of analyzed variables. Categorical variables were analyzed using the chi-square test, and results were reported as odds ratios with 95%CI. A Pearson's correlation analysis was used for an association between continuous parameters (age, cyst size, bilirubin level, and parity). A value less than 0.05 was considered statistically significant. Statistical analysis was performed via the SPSS version 26 (SPSS, Chicago, IL, United States).

## **RESULTS**

### *Study characteristics*

The PRISMA flow diagram (Figure 1) shows a total of 153 screened articles, while 71 articles met the inclusion criteria with a total of 97 patients. The earliest report was published in 1932, and the most recent was in 2020. Half of the studies were published in the last 2 decades (Figure 2). Most cases were from Asia ( $n = 35$ ), followed by Europe ( $n = 23$ ) and North America (Table 1). There were 10 non-English articles: two in Chinese, two in Japanese, one in Korean, 3 in German, one in French, and one in Spanish. Complete patient, cyst, and treatment characteristics were described in 55% of cases. All cases reported the mother's age, maternal outcome, and gestational age at delivery (Figure 3). Type of delivery was reported in 91%, and fetal outcome in 21.6% of cases.

### *Patient characteristics*

Patient characteristics are summarized in Table 2. The mean age was 25.3 years (range 15-40), and 62/83 (74.6%) patients with reported data were primigravidas. Parity was reported in 83 cases-62 women (74.6) were nulliparous, and seven had more than one



previous pregnancy. Among all, eight (8.2%) were diagnosed in the first, 33 (34.8%) in the second, 40 (41.2%) in the third trimester, and nine (9.3%) in the postpartum period. In six (6.1%) cases, CC was known before conception (Caroli disease).

### *Symptomatology and diagnostics*

The most common symptom was epigastric/right upper quadrant pain in 78 (80%) patients (Table 3). Jaundice was present in 58 (59%) and fever in 30 (30.1%). CC triad was described in 41 (50.5%) and the Charcot triad in 28 (28.8%) patients. Eight patients were asymptomatic, and six of them were diagnosed with Caroli disease before pregnancy. Most CCs were diagnosed using ultrasound ( $n = 58$ ) followed by MRI ( $n = 18$ ) and CT ( $n = 2$ ). In two cases, endoscopic retrograde cholangiopancreatography (ERCP) was employed as a diagnostic and therapeutic tool. Data were lacking for the rest of the patients, or the diagnosis was established intraoperatively.

In 28 patients (28.8%), imaging did not show dilation of intrahepatic ducts, while 22 (22.6%) had dilation. One patient had changes consistent with primary sclerosing cholangitis (PSC), while data on bile duct morphology were missing for 47 patients (48.4%).

Due to missing data, only bilirubin level was included in the analyses. It was reported in 70 cases (72.1%). The level was  $83.2 \pm 80.4$  mmol/L (range 5-426).

### *Cyst characteristics*

According to Todani's classification, cyst type was mentioned in 79 (81.4%) cases. The type was defined from published radiologic images in an additional five cases. Out of 84 cases, 56 (66.6%) patients were type I, 15 (17.8%) type IV, and 6 (7.1%) type V. In one case, the cyst was described as "transitory type", not included in Todani's classification. It may indicate cyst dilation of the CBD, which resolved after biliary drainage. The cyst size was described in 67 (69%) cases with a mean size of 12.5 cm (range 1.8-40). The cyst was ruptured at presentation in 6 cases during pregnancy and 1 case postpartum. Two

patients with ruptured CC died, resulting in a mortality rate of ruptured CC in pregnancy of 33%.

### ***Obstetric and surgical management***

The treatment approach was divided into four categories (Table 4), according to the management of CC and the management of pregnancy (Figure 4). Among those with radiologic, endoscopic, or surgical interventions, the main indication was cholangitis, followed by cyst rupture and progressive jaundice (Table 5 and Figure 5). The indication for urgent CC surgery was cyst rupture or CC-related bleeding in 11 cases. Outside these indications, the reasons for cesarean section (CS) were severe anemia (2%) and intense abdominal pain (3%). In seven cases, no signs of fetal compromise were present, but CS was performed to prevent potential complications of CC. In 29 (29.9%) patients, the treatment was expectant for both the cyst and pregnancy. These patients were primarily treated with antibiotics, bed rest, and intravenous fluids as a bridge therapy to the anticipated labor. Urgent CS was done in 24 (25%) patients, with indications shown in Table 5. Intervention for both the CC and pregnancy was done in 16 patients, and in four patients, cyst drainage/resection was performed simultaneously with CS.

### ***Maternal and fetal outcome***

The maternal outcome was mentioned in all cases. In ninety patients (91.7%), recovery after delivery and operation was uneventful, while seven (7.2%) died during or after pregnancy (Table 6).

Three cases of maternal death were published before 1950, and all were diagnosed during surgery for CC rupture/peritonitis. Martínez-Ordaz reported two patients who died postoperatively. The first patient died of postoperative bleeding at 9 wk' gestation following emergent surgery (HJ) for giant 20 cm CC with cholangitis. The second patient died after upper digestive tract bleeding and ischaemic perforation in the gastric fundus and hemoperitoneum following elective Roux-en-Y HJ made 4 wk postpartum. The cause of death was not defined. The third patient, in the 7<sup>th</sup> month of pregnancy, was

successfully treated for CC rupture with surgical drainage and ERCP. Still, the patient died of "some unknown cause". Another patient underwent postpartum hepaticoduodenostomy (HD) and died after 1 year from peritonitis of unknown cause (excluded from the mortality because the period is outside of puerperium). The fifth patient, at 20 wk' gestation, with typical CC symptoms and imminent abortion, underwent cyst resection and reconstruction and died from multiorgan failure and cholangitis postoperatively. In 1932, Zininger reported a patient at 4 mo gestation with cholangitis and CC diagnosed intraoperatively. The cyst was emptied, the anterior wall of the cyst was excised, and the defect was closed with sutures. The patient died eleven days after the operation from multiorgan failure due to progressive sepsis.

No complications related to labor were reported in patients who received interventional treatment of CC with the continuation of pregnancy. Regarding the postpartum treatment of patients who did not have definitive CC treatment during pregnancy ( $n = 76, 78.3\%$ ), surgical management was reported in 54 (71.0%), conservative in 8 (10.5%), while in 14 (18.4%) cases, further postpartum treatment was not specified. In surgically treated patients, after delivery, Roux-en-Y HJ was done in most cases (90.7%), followed by cystogastrostomy (5.5%), and cystoduodenostomy (3.7%). The average gestational age of pregnancy termination (induced and spontaneous) was 33.6 wk (range 9-40). Fetal outcomes were reported in 21 (21.6%) cases. However, we presumed that the authors would mention fetal or newborn death, so above mentioned fetal mortality could be reliable. From described cases, four children died postpartum, and two were stillborn. There were two spontaneous abortions (in the 8<sup>th</sup> and 20<sup>th</sup> gestational weeks) and two elective abortions (in the 20<sup>th</sup> and 14<sup>th</sup> weeks) due to cholangitis and progressive septic complications.

### *Correlation analyses*

Both cyst dimension and bilirubin level were described in 51 patients with a weak positive, statistically non-significant linear relationship ( $r = 0.32, P < 0.001$ ) (Figure 6). We found no relationship between age and cyst dimension ( $r = 0.06, P = 0.7$ ) for 67 patients

with available data. Similarly, no correlation was found between gestational age at cyst discovery and cyst size ( $r = 0.11$ ,  $P = 0.54$ ).

#### ***Univariate analyses of the factors related to the choice of treatment***

Cyst size is not necessarily related to the severity of symptoms, but cysts larger than 15 cm had a significantly higher likelihood of treatment intervention during pregnancy. Similarly, the Charcot triad was associated with a high likelihood of CC intervention (OR 4.3;  $P = 0.006$ ), induced labor (OR 5.4;  $P = 0.028$ ), and CC triad (OR 2.88;  $P = 0.017$ ). Other analyzed factors were not associated with the odds of interventional treatment (Table 7).

### **DISCUSSION**

#### ***Incidence***

Our study confirms the extreme rarity of maternal CC during pregnancy since less than 100 cases were collected. However, many cases of CC in pregnancy may not have been recognized or were asymptomatic, so the incidence is probably underestimated. Our collection of patients is the largest so far, and this study will contribute to a more precise estimation of the incidence of CC in pregnancy. Articles were systematically screened and reviewed, and in only two cases, we could not extract necessary data since we did not get a response from one author who presented a case in a Slovak journal<sup>[19]</sup>, and we could not adequately translate a case from Alizadeh *et al*<sup>[20]</sup> due to the Arabic alphabet and language.

#### ***Patient and disease characteristics***

Similarities and differences between CC in the general and pregnant populations exist. First, the distribution of CC types appears similar. In pregnancy, type I is the most common (73.8%), followed by type IV, comparable to the general population, where type I ranges between 80%-90%. Types II and III have not been reported, while Caroli disease occurred in six cases, corresponding to an incidence of 7.1%, several times higher than in the general population (1%-2%)<sup>[2,3,21]</sup>. Four cases of Caroli disease were managed

expectantly, and elective CS was performed in two cases, indicating a better prognosis than in newly diagnosed symptomatic CCs type I or IV. However, a better prognosis of Caroli disease in pregnancy may be attributed to a milder clinical course than other CC types.

Second, the CC size differed compared to the general population. In our study, more than two-thirds of CC could be classified as giant since the largest diameter was > 10 cm. The mean reported size in the general population is 3-4 cm, rarely exceeding 9 cm<sup>[4,11,15]</sup>. Hormonal changes and the compressive effect of the gravid uterus on CC with consequent cyst enlargement could be causative. To confirm this, serial sonographic exams and measurements of CC size in pregnant women in future studies are mandatory. Pregnancy may trigger CC symptoms in previously asymptomatic cases due to increasing CC size, which, along with physiological changes during pregnancy, may precipitate symptoms related to CC, such as jaundice, abdominal pain, or abdominal mass. This hypothesis is supported by the fact that almost 91% of CC from our study were discovered in the 2<sup>nd</sup> or 3<sup>rd</sup> trimester when the gravid uterus reaches the upper abdomen<sup>[9]</sup>. There is no data on the incidence of CC triad in the general population for females. However, a typical CC triad in more than 40% of women in our study indicates that the CC triad could be severalfold higher in pregnant women than in the general population (10%-20%)<sup>[11,12]</sup>. The incidence of CC rupture in our study was 7/97 (7.1 %),  
18 which is higher than in the general population (1%-2%)<sup>[10]</sup>. Cyst rupture was seen in 6 cases during pregnancy and one postpartum. Most (5/7) were reported in studies before 1990. Today, the wide availability of ultrasound, more frequent biochemical laboratory testing, and closer monitoring of pregnant women should maximally reduce the rate of severe consequences of CC. No CC rupture was noted during spontaneous or induced vaginal labor. Therefore, it remains debatable if vaginal delivery increases the risk of CC rupture.

Nausea and vomiting were reported in 37.5%, comparable to previous studies<sup>[6,11,22]</sup>. Interpretation of these symptoms is difficult as it is a common consequence of hormonal changes during pregnancy. CC resulting in duodenal or pyloric compression was an

indication for surgery in 3 cases. Abdominal distension or mass may be more prominent than in nonpregnant patients with CC. However, the women may neglect this symptom as it may be attributed to a normal pregnancy. The pain in the upper abdomen or right subcostal area should result in evaluating potential causes other than physiological pressure from the growing uterus. ICP, acute fatty liver of pregnancy, or other cholestatic disorders may contribute to the complexity of CC diagnosis and treatment (3% of pregnant women are affected by liver disorders during pregnancy<sup>[16,23]</sup>). Hence, other liver and biliary diseases, especially cholestatic, may lead to earlier or more pronounced jaundice in CC patients or could be the only cause of cholestasis unrelated to an asymptomatic CC pregnant patient.

### ***Treatment***

Due to the risk of malignancy, complete excision with biliodigestive reconstruction is the procedure of choice for nonpregnant patients with type I, II, or IV CC. Type III cysts may be managed with ERCP or endoscopic resection<sup>[5,24,25]</sup>. Our results show that Roux-en-Y HJ was a predominant operation in patients requiring cyst resection and reconstruction (92%). In two cases, HD was reported, and hepaticogastrostomy in one<sup>[26-28]</sup>.

The variety of reported treatment approaches indicates no standardized management of such a clinical scenario. Treatment was based on previous knowledge and experience related to CC in nonpregnant adults and general obstetric principles. The application of novel and minimally invasive percutaneous or endoscopic methods depended on the institutional availability of these procedures and the year of publication. Since this is a retrospective analysis, no control groups were available, and reliable comparative analysis of treatment approaches was not possible. However, by analyzing the severity of the clinical presentation, chosen medical approach, and outcomes, we constructed an algorithm that may help to guide clinical decisions (Figure 7).

The treatment in asymptomatic patients or smaller CC without cholangitis or gastric outlet obstruction includes observation until term, with planned postpartum CC management. Almost half cases (47.3%) did not require any intervention for CC. Even

when the labor was induced, in most cases, it was indicated more as a precaution than urgent obstetric indication, reported in only 39%. A prerequisite for conservative treatment is the absence of surgical or obstetric urgencies. It includes antibiotic therapy for septic complications, hospitalization, and more frequent follow-ups after discharge to prevent delayed identification of fetal compromise or CC rupture.

The decisive clinical characteristics that guided the treatment approach were the presence of cholangitis, cyst size, and to a lower degree, the presence of CC triad (Table 8). Parity, the mother's age, or jaundice alone did not affect the treatment choice. In cases of symptomatic CC, a conservative approach should be initiated, comprising analgesia, antibiotics, dietary changes, and ursodeoxycholic acid (in Caroli disease). Further steps depend on the course of the disease.

In only 22.9% of cases, open surgery, cyst excision, and reconstruction were done during pregnancy. The majority (81.0%) were operated on in the first trimester (54.5%). Possibly, most authors adopted a more conservative approach, particularly in the 2<sup>nd</sup> and 3<sup>rd</sup> trimesters. This is due to: (1) the minimization of the risk of complications from major surgery; and (2) the availability of less invasive methods for preventing CC complications.

In patients who require intervention, the minimally invasive methods have a high success rate. Only 3/22 (13.6%) cases required surgery for CC complications after unsuccessful percutaneous drainage/ERCP. Percutaneous cyst drainage may resolve symptoms, decrease the risk of cholangitis, and allow further intrauterine fetal development until viable gestational age<sup>[8,29,30]</sup>. On the other side, such procedures may carry a risk of pregnancy complications, although no severe complications of percutaneous drainage were reported in the present study. It remains unclear whether endoscopic or radiologic drainage should be performed for asymptomatic or mildly symptomatic cysts during the 3<sup>rd</sup> trimester since such procedures may trigger premature labor, while commonly, symptoms resolve on conservative measures.

Surgical urgencies (cyst rupture/peritonitis, massive bleeding, or severe cholangitis) mandate immediate "damage control" procedures such as internal/external cyst



drainage, T-drainage, or cholecystostomy. Synchronous CS and cyst excision plus HJ were done in only two cases in the earlier study period. Therefore, synchronous surgical and obstetric procedures should be done exceptionally.

In conservatively treated women, the timing of surgery after the delivery was 6-8 wk postpartum in most cases. No cases of CC complications during this "waiting" period were reported, so we consider this to be a sufficient recovery time for the patient without increased CC complications related to delayed management.

The treatment of Caroli disease generally differs from other CC types in the general population, and the same principles can be applied to Caroli's disease in pregnancy<sup>[31-33]</sup>. It includes supportive care with antibiotics for cholangitis and ursodeoxycholic acid for hepatolithiasis<sup>[31,34,35]</sup>. Surgical resection is considered with monolobar disease, but rarely as an urgent procedure<sup>[36]</sup>. The ursodeoxycholic acid may improve bile flow and reduce the incidence of biliary stones and cholangitis. It has been used to treat cholangiopathies, including primary biliary cholangitis or PSC. Studies suggest that ursodeoxycholic acid is effective and safe in patients with ICP<sup>[37]</sup>. However, its benefits in CC types 1-4 should yet be evaluated.

### ***Diagnostics***

Before the wide clinical usage of transabdominal ultrasound, CC during pregnancy was diagnosed intraoperatively. Since 1976, radiologic methods have enabled preoperative CC diagnosis. MRI became the common diagnostic method for pregnancy disorders in developed countries after 2000<sup>[38,39]</sup>. Consequently, cases from the first half of the 20th century were mostly diagnosed intraoperatively or at autopsy. Although emergent MRI is still not widespread, we strongly recommend MRI for newly sonographically detected CCs, as MRI defines the exact size and location of the cyst, stones, concomitant biliary anomalies, and compressive effect on surrounding organs, including the uterus.

### ***Maternal mortality***

The maternal mortality from CC was 7.2 %, but the accurate mortality appears lower since, in only 4 cases, a direct association with CC was confirmed. Publication bias (not reporting unfavorable outcomes) may have affected final mortality rates. In addition, in 5/7 fatal cases, important variables (bilirubin levels and cyst size) were not reported. Therefore, inferential analysis related to predictive factors for maternal mortality was not appropriate in this setting, particularly if we consider the study design that covered a long period (1932-2020) which is the main reason for heterogeneous diagnostic and therapeutic approaches. Clearly, in earlier studies, the diagnosis was commonly made intraoperatively. Radiologic, endoscopic, and intensive care procedures improved significantly over time<sup>[35,40,41]</sup>. Consequently, fetal and maternal mortality rates and complications are incomparable to results from more recent studies. Instead, we presented shortly every single case of maternal mortality in the results section and used multivariate analysis of factors that contributed to treatment choice.

### *Limitations of the study*

There are several limitations of the study. The first is missing data. In many cases, laboratory data and fetal outcome information are sparse. Indications for CS were not mentioned in many cases, as well as information on clinical presentation and cyst pathology. However, the most important CC characteristics and maternal outcomes were reported at acceptable rates. The second limitation was the inability to collect all cases due to language or unavailable full text. Third, a more thorough comparative statistical analysis could not be conducted regarding the design of the study, long study period, lack of data, and different treatment standards.

### **CONCLUSION**

CC in pregnancy presents a great therapeutic dilemma and an under-represented topic in the literature. Our results might contribute to understanding the behavior of CC in pregnancy. Although maternal death was rare, it can occur and was reported predominantly in studies before 1990. Without cholangitis or urgent surgical CC

complications, premature labor and fetal loss are uncommon. *Treatment should be tailored* to the individual patient and primarily based on gestational age, the risk of septic complications, and CC rupture. Careful monitoring and early recognition of CC complications are crucial for favorable maternal and fetal outcomes. CS is indicated when fetal distress, surgical emergencies, progressive cholangitis, or refractory symptoms related to CC exist. In other cases, conservative or minimally invasive CC treatment and an expectant obstetric approach should be the preferred initial approach.

## **ARTICLE HIGHLIGHTS**

### ***Research background***

No systematic data on choledochal cysts (CC) in pregnancy.

### ***Research motivation***

Due to the rarity, no guidelines exist for the diagnostic workup and treatment and obstetric strategy for CC in pregnancy.

### ***Research objectives***

To collect the most published case reports on CC in pregnancy.

### ***Research methods***

Descriptive statistics of available patient and disease data.

### ***Research results***

Cholangitis, CC > 15 cm, and bilirubin levels (> 80 mmol/L) were associated with a higher likelihood of urgent cesarean section (CS) and surgical intervention for CC. Bilirubin levels positively correlated with CC size. There was no correlation between age and cyst dimension, gestational age at cyst discovery, and CC size.

### ***Research conclusions***

Although rare, maternal CC in pregnancy should be included in the evaluation of jaundice with upper abdominal pain. Symptomatology and clinical course are variable. Treatment may range from an expectative approach to emergent surgical CC treatment and urgent CS. While most cases were managed by conservative measures or drainage procedures, CC > 15 cm and progressive cholangitis carry the risk of CC rupture and septic complications, which may increase the rates of unfavorable maternal and fetal outcomes.

### *Research perspectives*

All cases of CC in pregnancy should be published to understand better the epidemiology, etiology, specific diagnostic imaging methods, and treatment strategy of this extremely rare condition.

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SIMILARITY INDEX

### PRIMARY SOURCES

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