

# World Journal of *Gastrointestinal Endoscopy*

Submit a Manuscript: http://www.wjgnet.com/esps/ Help Desk: http://www.wjgnet.com/esps/helpdesk.aspx DOI: 10.4253/wjge.v7.i2.128 World J Gastrointest Endosc 2015 February 16; 7(2): 128-134 ISSN 1948-5190 (online) © 2015 Baishideng Publishing Group Inc. All rights reserved.

ORIGINAL ARTICLE

#### **Retrospective Study**

## Endoscopic retrograde cholangiopancreatography for suspected choledocholithiasis: From guidelines to clinical practice

#### Joana Magalhães, Bruno Rosa, José Cotter

Joana Magalhães, Bruno Rosa, José Cotter, Gastroenterology Department, Centro Hospitalar do Alto Ave, 4835-044 Guimarães, Portugal

José Cotter, Life and Health Sciences Research Institute (ICVS), School of Health Sciences, University of Minho, 4710-057 Braga, Guimarães, Portugal

José Cotter, ICVS/3B's, PT Government Associate Laboratory, 4710-057 Braga, Guimarães, Portugal

Author contributions: Magalhães J participated in the design of the study, performed data analysis and literature research and drafted the manuscript; Rosa B performed literature research and critically revised the manuscript; Cotter J critically revised the manuscript and approved the final version to be submitted; all the authors read and approved the final manuscript;

Ethics approval: This study was approved by the institutional review board of Centro Hospitalar do Alto Ave, Guimarães, Portugal.

**Informed consent:** All patients provided written consent to undergo endoscopic retrograde cholangiopancreatography and were informed of the risks and potential benefits of the procedure. **Conflict-of-interest:** The authors declare that there is no conflict of interests regarding the publication of this paper.

Data sharing: Technical appendix, statistical code, and dataset available from the corresponding author at jmagalhaes@chaa. min-saude.pt. The consent of the participants was not obtained but the presented data are anonymized and risk of identification is low.

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

Correspondence to: Joana Magalhães, MD, Gastroenterology Department, Centro Hospitalar do Alto Ave, Rua dos Cutileiros, Creixomil, 4835-044 Guimarães,

Portugal. jmagalhaes@chaa.min-saude.pt Telephone: +351-253-540330 Fax: +351-253-421308 Received: September 17, 2014 Peer-review started: September 20, 2014 First decision: October 14, 2014 Revised: December 14, 2014 Accepted: December 29, 2014 Article in press: December 31, 2014 Published online: February 16, 2015

## Abstract

**AIM:** To study the practical applicability of the American Society for Gastrointestinal Endoscopy guidelines in suspected cases of choledocholithiasis.

METHODS: This was a retrospective single center study, covering a 4-year period, from January 2010 to December 2013. All patients who underwent endoscopic retrograde cholangiopancreatography (ERCP) for suspected choledocholithiasis were included. Based on the presence or absence of predictors of choledocholithiasis (clinical ascending cholangitis, common bile duct (CBD) stones on ultrasonography (US), total bilirubin > 4 mg/dL, dilated CBD on US, total bilirubin 1.8-4 mg/dL, abnormal liver function test, age > 55 years and gallstone pancreatitis), patients were stratified in low, intermediate or high risk for choledocholithiasis. For each predictor and risk group we used the  $\chi^2$  to evaluate the statistical associations with the presence of choledocolithiasis at ERCP. Statistical analysis was performed using SPSS version 21.0. A P value of less than 0.05 was considered statistically significant.

**RESULTS:** A total of 268 ERCPs were performed for suspected choledocholithiasis. Except for gallstone pancreatitis (P = 0.063), all other predictors of cho-



ledocholitiasis (clinical ascending cholangitis, P = 0.001; CBD stones on US,  $P \leq 0.001$ ; total bilirubin > 4 mg/ dL, P = 0.035; total bilirubin 1.8-4 mg/dL, P = 0.001; dilated CBD on US,  $P \leq 0.001$ ; abnormal liver function test, P = 0.012; age > 55 years, P = 0.002) showed a statistically significant association with the presence of choledocholithiasis at ERCP. Approximately four fifths of patients in the high risk group (79.8%, 154/193 patients) had confirmed choledocholithiasis on ERCP, *vs* 34.2% (25/73 patients) and 0 (0/2 patients) in the intermediate and low risk groups, respectively. The definition of "high risk group" had a sensitivity of 86%, positive predictive value 79.8% and specificity 56.2% for the presence of choledocholithiasis at ERCP.

**CONCLUSION:** The guidelines should be considered to optimize patients' selection for ERCP. For high risk patients specificity is still low, meaning that some patients perform ERCP unnecessarily.

**Key words:** Choledocholithiasis; Endoscopic retrograde cholangiopancreatography; Cholangitis; Common bile duct stones; Dilated common bile duct

© **The Author(s) 2015.** Published by Baishideng Publishing Group Inc. All rights reserved.

**Core tip:** The American Society for Gastrointestinal Endoscopy (ASGE) proposes a stratification of patients according to the risk for choledocholithiasis, influencing subsequent management. Our study shown that the risk stratification, according to ASGE guidelines, may improve risk estimation of choledocholithiasis and should be considered to optimize patients' selection for endoscopic retrograde cholangiopancreatography (ERCP). However, even in the "high risk group" the specificity was low. Thus, at this point, it seems advisable that also "high risk" patients undergo further testing before being submitted to ERCP, similarly to those patients with "intermediate risk", while for patients with "low-risk" of choledocholithiasis a watchful waiting strategy seems adequate.

Magalhães J, Rosa B, Cotter J. Endoscopic retrograde cholangiopancreatography for suspected choledocholithiasis: From guidelines to clinical practice. *World J Gastrointest Endosc* 2015; 7(2): 128-134 Available from: URL: http://www. wjgnet.com/1948-5190/full/v7/i2/128.htm DOI: http://dx.doi. org/10.4253/wjge.v7.i2.128

## INTRODUCTION

Choledocholithiasis is the most common cause of biliary obstruction. Approximately 5% to 22% of the Western population has gallstones<sup>[1]</sup> and common bile duct stones occur in 8%-20%<sup>[2,3]</sup> of those patients. Patients suspected of having choledocholithiasis are

diagnosed with a combination of laboratory tests and imaging studies<sup>[4]</sup>. The first imaging study obtained is typically a transabdominal ultrasonography (US). When the ultrasound findings are not enough for a diagnosis a magnetic resonance cholangiopancreatography (MRCP) or an endoscopic ultrasound (EUS) should be considered.

The diagnosis of choledocholithiasis usually should be followed by some therapeutic intervention to remove the stones<sup>[4-7]</sup>. Endoscopic retrograde cholangiopancreatography (ERCP) is the standard method for the diagnosis and therapy of bile duct stones, however it is an invasive procedure not free of complications<sup>[8-11]</sup>.

According to the results of laboratory tests and US, the American Society for Gastrointestinal Endoscopy (ASGE) proposes to stratify a patient in low, intermediate or high risk for choledocholithiasis. Subsequent management will vary depending on the patient's level of risk<sup>[12]</sup>. The purpose of this study was to evaluate the practical applicability of the American Society for Gastrointestinal Endoscopy guidelines in a population of patients undergoing ERCP for suspected choledocholithiasis.

## MATERIALS AND METHODS

#### Patients

We performed a retrospective single center study, covering a 4-year period, from January 2010 to December 2013. Patients referred for ERCP for suspected bile duct lithiasis were consecutively included. Patients presenting for stent exchange or follow-up of known and incompletely removed stones on previous ERCP were excluded.

Clinical data recorded from disease onset (age, gender, symptoms at presentation, laboratorial values) to the time of the ERCP (therapeutic procedures and related complications) were collected.

### Predictors of choledocholithiasis

According to ASGE guidelines<sup>[12]</sup>, cholangitis, total bilirubin > 4 mg/dL and common bile duct (CBD) stone on US were considered very strong predictors. Total bilirubin 1.8-4 mg/dL and dilated CBD on US were considered strong predictors and abnormal liver biochemical tests, age > 55 years and gallstone pancreatitis were considered moderate predictors. Patients with strong predictors or any very strong predictor were considered at high risk for choledocholithiasis. Patients without any predictor and all other patients were considered low and intermediate risk for choledocholithiasis, respectively. The diagnosis of cholangitis was established by the presence of Charcot's triad (fever, abdominal pain and jaundice). The diagnosis of CBD stone on US was considered when an intraductal echogenic focus with distal acoustic

Baishideng®

Magalhães J et al. Endoscopic retrograde cholangiopancreatography for suspected choledocholithiasis

Table 1	Baseline cl	haracteristics of	the stud	ly populatio	1 <i>n</i> (%)
---------	-------------	-------------------	----------	--------------	----------------

Variable	Total ( $n = 268$ )
Gender, female	161 (60.1)
Age, mean ± SD	$66.8 \pm 16.8$
Very strong predictors	
Clinical ascending cholangitis	36 (13.4)
Common bile duct stone on US	109 (40.7)
Total bilirubin > 4 mg/dL	102 (38.1)
Strong predictors	
Total bilirubin 1.8-4 mg/dL	84 (31.3)
Dilated common bile duct on US	195 (72.8)
Moderate predictors	
Abnormal liver function test	231 (86.2)
Age > 55 yr	197 (73.5)
Gallstone pancreatitis	63 (23.5)

US: Ultrasonography.

shadow was identified. Dilated CBD on US was considered when bile duct diameter was > 6 mm in a patient without cholecystectomy. Abnormal liver biochemical tests were considered when aspartate aminotransferase (AST), alanine aminotransferase (ALT) and alkaline phosphatase (AP) presented elevated laboratory values, considering the reference lab values in our institution. Gallstone pancreatitis was considered when patients presented with abdominal pain (epigastric pain often radiating to the back), lipase (or amylase activity) at least 3 times higher than the upper limit of normal, stones or biliary sludge within gallbladder and no history of alcohol abuse.

## Endoscopic retrograde cholangiopancreatography procedure

Every ERCP was performed using Olympus<sup>®</sup> TJF 160 VR or TJF 145 side-viewing endoscopes. All patients provided written consent to undergo ERCP and were informed of the risks and benefits of the procedure. Patients were under propofol sedation assisted by an anaesthesiologist. Stone size and number were documented on the initial diagnostic cholangiogram at ERCP. Endoscopic sphincterotomy was performed over a guide wire. Some patients underwent papillary balloon dilation using a through-the-scope balloon catheter for oesophageal/pyloric dilation, gradually inflated to 12-18 mm according to the size of the largest stone and the maximal diameter of the distal bile duct on the cholangiogram. Stones were removed using a retrieval balloon catheter and/or a Dormia basket. When necessary, mechanical lithotripsy was performed to fragment the stones prior to removal. Complete clearance of the bile duct was documented with a balloon catheter cholangiogram at the end of the procedure. In the case of residual lithiasis, a biliary 7 Fr double pigtail plastic stent was placed and a second ERCP was planned within 10-12 wk. At the end of each ERCP, 100 mg rectal indomethacin was routinely given, to prevent post-ERCP pancreatitis. Prophylactic antibiotics were not routinely administered.

#### Statistical analysis

Statistical analysis was performed using SPSS version 21.0 (SPSS<sup>®</sup> Inc., Chicago, IL, United States).

Quantitative data were described as mean  $\pm$  SD and qualitative data as proportions. For each predictor and risk group the  $\chi^2$  was used to access differences between presence *vs* absence of choledocolithiasis on ERCP. A *P* value < 0.05 was considered statistically significant.

For each risk group and their predictors the sensitivity, specificity, positive predictive values (PPV) and negative predictive value (NPV) were assessed.

#### RESULTS

From January 2010 to December 2013, a total of 268 patients were referred for ERCP for suspected choledocholithiasis. Patients included in our study were predominantly female (60.1%), with a mean age of  $66.8 \pm 16.8$  years. Choledocholithiasis was present in 179 ERCPs (66.8%). The predictors more often seen in our patients were the presence of abnormal liver biochemical tests (86.2%), age > 55 years (73.5%) and dilated CBD on US (72.8%). Main clinical features of the study population are shown in Table 1.

#### Predictors of choledocholithiasis

Except for gallstone pancreatitis (P = 0.063), all other predictors showed a statistically significant difference between presence *vs* absence of choledocholithiasis on ERCP (cholangitis, P = 0.001; CBD stone on US, P < 0.001; total bilirubin > 4 mg/dL, P = 0.035; total bilirubin 1.8-4 mg/dL, P = 0.001; dilated CBD on US, P < 0.001; abnormal liver function test, P = 0.012; age > 55 years, P = 0.002) (Table 2).

The risk of choledocholithiasis, as shown by *odds ratio*, was increased for patients who presented with cholangitis (OR: 6.48, 95%CI: 1.93-21.80), common bile duct stone on US (OR: 11.25, 95%CI: 5.32-23.81), total bilirubin > 4 mg/dL (OR: 1.79, 95%CI: 1.04-3.08), total bilirubin 1.8-4 mg/dL (OR:3.15, 95%CI: 1.63-6.08), dilated common bile duct on US (OR:5.06, 95%CI: 2.85-8.99), abnormal liver function test (OR:2.43, 95%CI: 1.20-4.90) and age > 55 years (OR:2.37, 95%CI: 1.36-4.15) (Table 2).

#### Risk group for choledocholithiasis

Of the 268 patients included in this study, 72% were stratified into the high risk group. Of the remaining patients, 27.2% e 0.8% were stratified into the intermediate and low risk groups, respectively. Approximately four fifths of patients in the high risk group (79.8%, 154/193 patients) had confirmed choledocholithiasis on ERCP. The presence of choledocholithiasis was identified in 34.2% (25/73) of intermediate risk patients. Any patient into the low risk group had choledocholithiasis on ERCP. There was a statistically significant association between



Table 2 Predictors of choledocholithiasis - univariate analysis n (%)					
Variable	Choledocholithiasis on ERCP	No Choledocholithiasis on ERCP	OR	95%CI	P value
Very strong predictors					
Clinical ascending cholangitis	33 (91.7)	3 (8.3)	6.48	1.93-21.80	0.001
Common bile duct stone on US	100 (91.7)	9 (8.3)	11.25	5.32-23.81	< 0.001
Total bilirubin > 4 mg/dL	76 (74.5)	26 (25.5)	1.79	1.04-3.08	0.035
Strong predictors					
Total bilirubin 1.8-4 mg/dL	63 (75.0)	21 (25.0)	3.15	1.63-6.08	0.001
Dilated common bile duct on US	150 (76.9)	45 (23.1)	5.06	2.85-8.99	< 0.001
Moderate predictors					
Abnormal liver function test	161 (69.7)	70 (30.3)	2.43	1.20-4.90	0.012
Age > 55 yr	142 (72.1)	55 (27.9)	2.37	1.36-4.15	0.002
Gallstone pancreatitis	36 (57.2)	27 (42.8)	0.58	0.32-1.03	0.063

ERCP: Endoscopic retrograde cholangiopancreatography; US: Ultrasonography.

Table 3 Risk group for choledocholithiasis - univariate analysis $n$ (%)						
Variable	Total	Choledocholithiasis on ERCP	No Choledocholithiasis on ERCP	<b>P</b> value		
High risk group	193 (72.0)	154 (79.8)	39 (20.2)	< 0.001		
Intermediate risk group	73 (27.2)	25 (34.2)	48 (65.8)			
Low risk group	2 (0.8)	0 (0)	2 (100)			
Very strong predictors						
None	97 (36.2)	39 (40.2)	58 (59.8)	< 0.001		
One	104 (38.8)	80 (76.9)	24 (23.1)			
Two	58 (21.6)	51 (87.9)	7 (12.1)			
Three	9 (3.4)	9 (100)	0 (0)			
Strong predictors						
None	27 (16.4)	3 (11.1)	24 (88.9)	< 0.001		
One	78 (47.3)	50 (64.1)	28 (35.9)			
Two	60 (36.4)	50 (83.3)	10 (16.7)			

ERCP: Endoscopic retrograde cholangiopancreatography.

the presence of choledocholithiasis on ERCP and the risk group (P < 0.001) (Table 3). The odds ratio (OR) for choledocholithiasis in high risk patients was 7.89 (95%CI: 4.36-14.32). The combination of any two or all very strong predictors elevated the probability of choledocholithiasis for 87.9% (51/58) and 100% (9/9), respectively. The combination of both strong predictors presented 83.3% (50/60) of probability of choledocolithiasis.

## Sensitivity, specificity, positive predictive values and negative predictive values for choledocolithiasis

Cholangitis was the parameter that had the higher specificity (96.6%), however for the same parameter the sensitivity was low. Total bilirubin > 4 mg/dL or the presence of CBD stones on US also presented a good specificity (89.9% and 70.8%, respectively). The PPV was high for very strong predictors, mainly clinical ascending cholangitis (PPV 91.7%) and CBD stones on US (PPV 91.7%). The high risk group had a high sensitivity (86%) and PPV (79.8%), but low specificity (56.2%) for the presence of CBD stones (Table 4).

#### DISCUSSION

According to ASGE guidelines, a patient stratified as

high risk has > 50% of probability of choledocholithiasis<sup>[12]</sup>. In our study, patients stratified as high risk following ASGE criteria had 79.8% probability of choledocholithiasis. These results are consistent with those presented in the study by Rubin *et al*<sup>[13]</sup>. All the very strong predictors (clinical ascending cholangitis, CBD stones on US or total bilirubin > 4 mg/dL) presented a statistically significant association with the presence of choledocholithiasis. The combination of any of two or three very strong predictors increased the probability of choledocholithiasis for 87.9% and 100%, respectively.

Transabdominal ultrasound is the most commonly used initial imaging modality for suspected biliary stones. In our study, the presence of CBD stones detected during the US evaluation presented an OR of 11.25 for choledocholithisis. The diagnosis of choledocholithiasis is often difficult, with the sensitivity for the detection of CBD stones by US ranging from 20% to 80%<sup>[14]</sup>. The diagnostic accuracy of US is operator dependent but it is also influenced by some clinical features of patients (shadowing from bowel gas, overweight and stone size)<sup>[14]</sup>.

In our study, the combination of strong predictors (dilated CBD on US, total bilirubin 1.8-4 mg/dL) presented 83.3% of probability of choledocholithiasis confirmed at ERCP. Strong predictors presented a sta-

Table 4   Sensitivity, specificity, positive predictive values and negative predictive values for choledocolithiasis					
Variable	Sensitivity	Specificity	PPV	NPV	
Very strong predictors					
Clinical ascending cholangitis	18.4	96.6	91.7	37.0	
Common bile duct stone on US	55.9	89.9	91.7	50.3	
Total bilirubin > 4 mg/dL	42.5	70.8	74.5	37.8	
Strong predictors					
Total bilirubin 1.8-4 mg/dL	61.1	66.6	75	51.2	
Dilated common bile duct on US	83.8	49.4	76.9	60.3	
Moderate predictors					
Abnormal liver function test	89.9	21.3	69.7	51.3	
Age > 55 yr	79.3	38.2	72.1	47.9	
Gallstone pancreatitis	20.1	69.7	57.1	30.2	
High risk group	86	56.2	79.8	66.7	
Intermediate risk group	13.9	46	34,2	21	

PPV: Positive predictive values; NPV: Negative predictive values; US: Ultrasonography.

tistically significant association with the presence of choledocholithiasis, which is in line with other previously published data<sup>[15-18]</sup>. The OR for choledocholithiasis in a patient with a CBD dilation was 5.06. However, the CBD dilation should always be interpreted according to patient characteristics, particularly previous cholecystectomy and age<sup>[19-21]</sup>. Previous studies<sup>[15-17,22,23]</sup> have reported some utility of serum bilirubin levels as a predictor of CBD stones. In this study, a bilirubin value between 1.8-4 g/dL had an OR of 3.15 and a specificity of 66.6% for choledocolithiasis. The specificity increased to 70% when the bilirrubin value was > 4 mg/dL. These results are in agreement with those previously reported by ASGE guidelines<sup>[12]</sup>.

Individually, moderate predictors, such as abnormal liver function test and age > 55 years, presented a statistically significant association with the presence of choledocholithiasis in our series and a sensitivity of 89.9% and 79.3% for the prediction of choledocholithiasis on ERCP. In a study by Barkun *et al*<sup>[16]</sup>, abnormal liver function tests, such as alkaline phosphatase > 300 units/L and AST > 120 units/L present a sensitivity of 79% and 81% to predict choledocholithiasis, respectively. At the same study, age > 55 years, only presented a sensitivity of 57%, however, when combined with other predictors (elevated bilirubin and CBD dilation on US) the model predicted with 94% of probability the presence of choledocholithiasis.

As previously reported by other authors<sup>[13,24]</sup>, also in our results the diagnosis of gallstone pancreatitis was not related with the presence of choledocholithiasis at ERCP (P > 0.05). Stone size may be an explanation, as larger stones are less likely to migrate<sup>[24]</sup> and the small gallstones, that most commonly are the source of pancreatitis<sup>[25]</sup>, frequently pass spontaneously. Some studies have reported that in the absence of cholangitis, patients with gallstone pancreatitis do not benefit from early ERCP<sup>[26,27]</sup>.

In patients stratified into the intermediate and low risk group, the probability of choledocholithiasis is 10%-50% and < 10%, respectively<sup>[12]</sup>. In this study, the probability of choledocholithiasis was 34.2% (25/73) and 0 (0/2) for intermediate and low risk groups, respectively. For these risk groups the sensitivity, specificity, PPV and NPV did not show values with clinical interest. In the intermediate risk group, ASGE guidelines<sup>[12]</sup> recommended less invasive options for detecting choledocholithiasis, such as MRCP or EUS. The two techniques showed a good sensitivity and specificity for choledocholithiasis<sup>[28,29]</sup>, so deciding which test should be performed first depends on various factors such as availability, cost, patient-related factors and the suspicion for a small stone. Because it is noninvasive, MRCP is the first test performed to look for CBD stones. However, for small CBD stones (< 5 mm) the sensitivity of MRCP is lower<sup>[30]</sup>, so, if the MRCP is negative, but the suspicion for a common bile duct stone remains moderate to high, EUS is an appropriate next step.

In conclusion, our study confirms that the combination of choledocolithiasis predictors, according to ASGE guidelines<sup>[12]</sup>, enables risk stratification of patients based on the likelihood for the presence of choledocholithiasis. However, for high risk patients the specificity was still low (56.2%), with 39 patients (20%) false positive, meaning that a significant proportion of patients will be submitted to ERCP unnecessarily. In the future, the inclusion of new predictors or different combinations of previous predictors will be essential to improve the classification of patients as high risk, obviating the need of other imaging tests (MRCP/EUS) before ERCP. However, at this point, it seems advisable that also "high risk" patients undergo further testing with MRCP and/or EUS before being submitted to ERCP, similarly to those patients with "intermediate risk", while for patients with "lowrisk" of choledocholithiasis a watchful waiting strategy seems adequate.

#### COMMENTS

#### Background

Patients suspected of having choledocholithiasis are diagnosed with a combination of laboratory tests and/or imaging studies. Endoscopic retrograde cholangiopancreatography (ERCP) has been established as the standard method for the management of bile duct stones, but it may be associated with substantial morbidity and mortality. In the evaluation of suspected choledocolithiasis, the American Society for Gastrointestinal Endoscopy (ASGE) proposes to stratify a patient as high risk, intermediate risk or low risk for having choledocholithiasis. Subsequent management will vary depending on the patient's level of risk.

#### **Research frontiers**

In this study, the authors aimed to assess the practical applicability and to validate the current ASGE guidelines in a population of patients undergoing ERCP for suspected choledocholithiasis.

#### Innovations and breakthroughs

The study confirms that the combination of choledocolithiasis predictors, according to ASGE guidelines may improve risk estimation of choledocholithiasis and should be considered to optimize patients' selection for ERCP. However, even in the "high risk group" the specificity was low (56.2%), meaning that a significant proportion of patients will still perform ERCP unnecessarily.



#### Applications

The results of this study suggest that the inclusion of new predictors of choledocholithisis or different combinations of previous predictors will be essential to improve the classification of patients as high risk, obviating the need of other imaging tests before endoscopic retrograde cholangiopancreatography. Thus, at this point, it seems advisable that also "high risk" patients undergo further testing before being submitted to ERCP, similarly to those patients with "intermediate risk", while for patients with "low-risk" of choledocholithiasis a watchful waiting strategy seems adequate.

#### Terminology

Choledocholithiasis is defined as the occurrence of stones in the bile duct and has a propensity for life-threatening complications such as cholangitis and acute pancreatitis. Endoscopic retrograde cholangiopancreatography is a technique that combines the use of endoscopy and fluoroscopy to diagnose and treat problems of the biliary or pancreatic ductal systems. It has evolved from a diagnostic procedure to an almost exclusively therapeutic technique.

#### Peer-review

Title and running title accurately reflects the topic and contents of the paper key words.

### REFERENCES

- Kaltenthaler E, Vergel YB, Chilcott J, Thomas S, Blakeborough T, Walters SJ, Bouchier H. A systematic review and economic evaluation of magnetic resonance cholangiopancreatography compared with diagnostic endoscopic retrograde cholangiopancreatography. *Health Technol Assess* 2004; 8: iii, 1-89 [PMID: 14982656]
- Ko CW, Lee SP. Epidemiology and natural history of common bile duct stones and prediction of disease. *Gastrointest Endosc* 2002; 56: S165-S169 [PMID: 12447261]
- 3 Tazuma S. Gallstone disease: Epidemiology, pathogenesis, and classification of biliary stones (common bile duct and intrahepatic). Best Pract Res Clin Gastroenterol 2006; 20: 1075-1083 [PMID: 17127189]
- 4 **Caddy GR**, Tham TC. Gallstone disease: Symptoms, diagnosis and endoscopic management of common bile duct stones. *Best Pract Res Clin Gastroenterol* 2006; **20**: 1085-1101 [PMID: 17127190]
- 5 Williams EJ, Green J, Beckingham I, Parks R, Martin D, Lombard M. Guidelines on the management of common bile duct stones (CBDS). *Gut* 2008; 57: 1004-1021 [PMID: 18321943]
- 6 Scientific Committee of the European Association for Endoscopic Surgery (E.A.E.S.). Diagnosis and treatment of common bile duct stones (CBDS). Results of a consensus development conference. Surg Endosc 1998; 12: 856-864 [PMID: 9602006]
- 7 Maple JT, Ikenberry SO, Anderson MA, Appalaneni V, Decker GA, Early D, Evans JA, Fanelli RD, Fisher D, Fisher L, Fukami N, Hwang JH, Jain R, Jue T, Khan K, Krinsky ML, Malpas P, Ben-Menachem T, Sharaf RN, Dominitz JA. The role of endoscopy in the management of choledocholithiasis. *Gastrointest Endosc* 2011; 74: 731-744 [PMID: 21951472]
- 8 Loperfido S, Angelini G, Benedetti G, Chilovi F, Costan F, De Berardinis F, De Bernardin M, Ederle A, Fina P, Fratton A. Major early complications from diagnostic and therapeutic ERCP: a prospective multicenter study. *Gastrointest Endosc* 1998; **48**: 1-10 [PMID: 9684657]
- 9 Masci E, Toti G, Mariani A, Curioni S, Lomazzi A, Dinelli M, Minoli G, Crosta C, Comin U, Fertitta A, Prada A, Passoni GR, Testoni PA. Complications of diagnostic and therapeutic ERCP: a prospective multicenter study. *Am J Gastroenterol* 2001; 96: 417-423 [PMID: 11232684]
- 10 Christensen M, Matzen P, Schulze S, Rosenberg J. Complications of ERCP: a prospective study. *Gastrointest Endosc* 2004; 60: 721-731 [PMID: 15557948]
- 11 Anderson MA, Fisher L, Jain R, Evans JA, Appalaneni V, Ben-Menachem T, Cash BD, Decker GA, Early DS, Fanelli RD, Fisher DA, Fukami N, Hwang JH, Ikenberry SO, Jue TL, Khan KM, Krinsky ML, Malpas PM, Maple JT, Sharaf RN, Shergill AK,

Dominitz JA. Complications of ERCP. *Gastrointest Endosc* 2012; **75**: 467-473 [PMID: 22341094]

- 12 Maple JT, Ben-Menachem T, Anderson MA, Appalaneni V, Banerjee S, Cash BD, Fisher L, Harrison ME, Fanelli RD, Fukami N, Ikenberry SO, Jain R, Khan K, Krinsky ML, Strohmeyer L, Dominitz JA. The role of endoscopy in the evaluation of suspected choledocholithiasis. *Gastrointest Endosc* 2010; **71**: 1-9 [PMID: 20105473]
- 13 Rubin MI, Thosani NC, Tanikella R, Wolf DS, Fallon MB, Lukens FJ. Endoscopic retrograde cholangiopancreatography for suspected choledocholithiasis: testing the current guidelines. *Dig Liver Dis* 2013; 45: 744-749 [PMID: 23540659]
- 14 Mandelia A, Gupta AK, Verma DK, Sharma S. The Value of Magnetic Resonance Cholangio-Pancreatography (MRCP) in the Detection of Choledocholithiasis. *J Clin Diagn Res* 2013; 7: 1941-1945 [PMID: 24179904]
- 15 Abboud PA, Malet PF, Berlin JA, Staroscik R, Cabana MD, Clarke JR, Shea JA, Schwartz JS, Williams SV. Predictors of common bile duct stones prior to cholecystectomy: a meta-analysis. *Gastrointest Endosc* 1996; 44: 450-455 [PMID: 8905367]
- 16 Barkun AN, Barkun JS, Fried GM, Ghitulescu G, Steinmetz O, Pham C, Meakins JL, Goresky CA. Useful predictors of bile duct stones in patients undergoing laparoscopic cholecystectomy. McGill Gallstone Treatment Group. *Ann Surg* 1994; 220: 32-39 [PMID: 7517657]
- 17 Yang MH, Chen TH, Wang SE, Tsai YF, Su CH, Wu CW, Lui WY, Shyr YM. Biochemical predictors for absence of common bile duct stones in patients undergoing laparoscopic cholecystectomy. *Surg Endosc* 2008; 22: 1620-1624 [PMID: 18000708]
- 18 Prat F, Meduri B, Ducot B, Chiche R, Salimbeni-Bartolini R, Pelletier G. Prediction of common bile duct stones by noninvasive tests. *Ann Surg* 1999; 229: 362-368 [PMID: 10077048]
- 19 Daradkeh S, Tarawneh E, Al-Hadidy A. Factors affecting common bile duct diameter. *Hepatogastroenterology* 2005; 52: 1659-1661 [PMID: 16334751]
- 20 Park SM, Kim WS, Bae IH, Kim JH, Ryu DH, Jang LC, Choi JW. Common bile duct dilatation after cholecystectomy: a one-year prospective study. *J Korean Surg Soc* 2012; 83: 97-101 [PMID: 22880184]
- 21 Kaim A, Steinke K, Frank M, Enriquez R, Kirsch E, Bongartz G, Steinbrich W. Diameter of the common bile duct in the elderly patient: measurement by ultrasound. *Eur Radiol* 1998; 8: 1413-1415 [PMID: 9853225]
- 22 **Onken JE**, Brazer SR, Eisen GM, Williams DM, Bouras EP, DeLong ER, Long TT, Pancotto FS, Rhodes DL, Cotton PB. Predicting the presence of choledocholithiasis in patients with symptomatic cholelithiasis. *Am J Gastroenterol* 1996; **91**: 762-767 [PMID: 8677945]
- 23 Peng WK, Sheikh Z, Paterson-Brown S, Nixon SJ. Role of liver function tests in predicting common bile duct stones in acute calculous cholecystitis. *Br J Surg* 2005; 92: 1241-1247 [PMID: 16078299]
- 24 Tranter SE, Thompson MH. Spontaneous passage of bile duct stones: frequency of occurrence and relation to clinical presentation. Ann R Coll Surg Engl 2003; 85: 174-177 [PMID: 12831489]
- 25 Venneman NG, Buskens E, Besselink MG, Stads S, Go PM, Bosscha K, van Berge-Henegouwen GP, van Erpecum KJ. Small gallstones are associated with increased risk of acute pancreatitis: potential benefits of prophylactic cholecystectomy? *Am J Gastroenterol* 2005; **100**: 2540-2550 [PMID: 16279912]
- 26 Chang L, Lo SK, Stabile BE, Lewis RJ, de Virgilio C. Gallstone pancreatitis: a prospective study on the incidence of cholangitis and clinical predictors of retained common bile duct stones. *Am J Gastroenterol* 1998; **93**: 527-531 [PMID: 9576442]
- 27 Sakai Y, Tsuyuguchi T, Ishihara T, Yukisawa S, Ohara T, Tsuboi M, Ooka Y, Kato K, Katsuura K, Kimura M, Takahashi M, Nemoto K, Miyazaki M, Yokosuka O. Is ERCP really necessary in case of suspected spontaneous passage of bile duct stones? *World J Gastroenterol* 2009; 15: 3283-3287 [PMID: 19598305]

Magalhães J et al. Endoscopic retrograde cholangiopancreatography for suspected choledocholithiasis

- 28 Verma D, Kapadia A, Eisen GM, Adler DG. EUS vs MRCP for detection of choledocholithiasis. *Gastrointest Endosc* 2006; 64: 248-254 [PMID: 16860077]
- Ledro-Cano D. Suspected choledocholithiasis: endoscopic ultrasound or magnetic resonance cholangio-pancreatography? A systematic review. *Eur J Gastroenterol Hepatol* 2007; 19: 1007-1011 [PMID: 18049172]
- 30 Kondo S, Isayama H, Akahane M, Toda N, Sasahira N, Nakai Y, Yamamoto N, Hirano K, Komatsu Y, Tada M, Yoshida H, Kawabe T, Ohtomo K, Omata M. Detection of common bile duct stones: comparison between endoscopic ultrasonography, magnetic resonance cholangiography, and helical-computed-tomographic cholangiography. *Eur J Radiol* 2005; 54: 271-275 [PMID: 15837409]

P- Reviewer: Lee CL, Muguruma N, Skok P, Yu B S- Editor: Ji FF L- Editor: A E- Editor: Zhang DN







## Published by Baishideng Publishing Group Inc

8226 Regency Drive, Pleasanton, CA 94588, USA Telephone: +1-925-223-8242 Fax: +1-925-223-8243 E-mail: bpgoffice@wjgnet.com Help Desk: http://www.wjgnet.com/esps/helpdesk.aspx http://www.wjgnet.com

