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# Laparoscopic cholecystectomy versus percutaneous catheter drainage for acute cholecystitis in high risk patients (CHOCOLATE): multicentre randomised clinical trial

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## ABSTRACT

### OBJECTIVE

To assess whether laparoscopic cholecystectomy is superior to percutaneous catheter drainage in high risk patients with acute calculous cholecystitis.

### DESIGN

Multicentre, randomised controlled, superiority trial.

### SETTING

11 hospitals in the Netherlands, February 2011 to January 2016.

### PARTICIPANTS

142 high risk patients with acute calculous cholecystitis were randomly allocated to laparoscopic cholecystectomy (n=66) or to percutaneous catheter drainage (n=68). High risk was defined as an acute physiological assessment and chronic health evaluation II (APACHE II) score of 7 or more.

### MAIN OUTCOME MEASURES

The primary endpoints were death within one year and the occurrence of major complications, defined as infectious and cardiopulmonary complications within one month, need for reintervention (surgical, radiological, or endoscopic that had to be related to acute cholecystitis) within one year, or recurrent biliary disease within one year.

### RESULTS

The trial was concluded early after a planned interim analysis. The rate of death did not differ between the laparoscopic cholecystectomy and percutaneous catheter drainage group (3% v 9%, P=0.27), but major complications occurred in eight of 66 patients

(12%) assigned to cholecystectomy and in 44 of 68 patients (65%) assigned to percutaneous drainage (risk ratio 0.19, 95% confidence interval 0.10 to 0.37; P<0.001). In the drainage group 45 patients (66%) required a reintervention compared with eight patients (12%) in the cholecystectomy group (P<0.001). Recurrent biliary disease occurred more often in the percutaneous drainage group (53% v 5%, P<0.001), and the median length of hospital stay was longer (9 days v 5 days, P<0.001).

### CONCLUSION

Laparoscopic cholecystectomy compared with percutaneous catheter drainage reduced the rate of major complications in high risk patients with acute cholecystitis.

### TRIAL REGISTRATION

Dutch Trial Register NTR2666.

### Introduction

Acute cholecystitis is a common indication for hospital admission and an increasing burden on the Western healthcare system. In the United States, the number of hospital admissions for acute cholecystitis increased by 44% during 1997-2012, from 149 661 to 215 995.<sup>1</sup>

In young, otherwise healthy patients early laparoscopic cholecystectomy is considered the treatment of choice for acute calculous cholecystitis.<sup>2</sup> In high risk patients the management of acute cholecystitis remains controversial. Cholecystectomy in these patients can lead to serious morbidity and mortality owing to reduced physiological reserve.<sup>3-6</sup> Therefore, imaging guided percutaneous catheter drainage is increasingly being performed as an alternative to early cholecystectomy. This minimally invasive radiological procedure resolves local and systemic inflammation without the risks of surgery. According to international guidelines, it is a valuable treatment in high risk patients and in those with moderate or severe cholecystitis.<sup>7</sup> A drawback of percutaneous catheter drainage, however, is that it is not a definitive treatment since the gallbladder is not removed. This may lead to recurrent cholecystitis, and other biliary complications with severe clinical effects.<sup>8,9</sup>

No randomised studies have compared laparoscopic cholecystectomy with percutaneous catheter drainage in patients with acute cholecystitis. It therefore remains

## WHAT IS ALREADY KNOWN ON THIS TOPIC

No randomised studies have compared laparoscopic cholecystectomy with percutaneous catheter drainage in patients with acute calculous cholecystitis. It therefore remains unclear which treatment should be preferred in terms of clinical and economical outcomes in high risk patients.

## WHAT THIS STUDY ADDS

This study provides strong evidence that laparoscopic cholecystectomy is superior to percutaneous catheter drainage in the treatment of high risk patients with acute calculous cholecystitis.

Cholecystectomy not only reduced the rate of major complications (ie, infectious and cardiopulmonary complications, or need for reintervention, or recurrent biliary disease), but also reduced utilisation of healthcare resources and costs by more than 30%.

unclear which treatment should be preferred in terms of clinical and economical outcomes. In daily practice, both cholecystectomy and percutaneous catheter drainage are performed according to the preference of the treating surgeon, gastroenterologist, or other clinicians.

We performed a nationwide randomised trial (CHOCOLATE) to assess whether laparoscopic cholecystectomy is superior to percutaneous catheter drainage in high risk patients with acute calculous cholecystitis.

## Methods

### Study design and participants

The CHOCOLATE study was designed as a multicentre, randomised controlled, superiority trial, and the protocol has been previously described.<sup>10</sup> Adults with acute calculous cholecystitis and a high surgical risk were enrolled in 11 teaching hospitals in the Netherlands. Acute cholecystitis was defined according to the Tokyo guidelines.<sup>11</sup> Risk assessment was based on the APACHE II (acute physiology assessment and chronic health evaluation II) severity of disease classification system.<sup>12</sup> High risk was defined as an APACHE II score of 7 or more. We chose this cut-off on the basis of systematic evaluation of several imaginary case scenarios by a multicentre, multidisciplinary expert panel of surgeons, gastroenterologists, and radiologists. Patients with an APACHE II score of 15 or more were excluded because the risk of mortality in these patients was deemed too high—that is, disease severity or comorbidity, or both, presented a strict contraindication to surgery. We also excluded patients with symptoms that lasted longer than seven days at time of first presentation, since these patients should undergo delayed cholecystectomy according to the Dutch treatment guidelines.<sup>13</sup> Other exclusion criteria were pregnancy, decompensated liver cirrhosis, admission to the intensive care unit at the time of cholecystitis diagnosis, and mental illness prohibiting informed consent.

The study was investigator initiated and conducted in accordance with the principles of the Declaration of Helsinki. The protocol was approved by the institutional review board of each participating centre. The safety and efficacy of the trial was monitored by a data safety monitoring board consisting of three independent, non-participating clinicians and an independent epidemiologist. All patients or their legal representatives provided written informed consent.

### Randomisation and masking

Patients were randomly assigned to either laparoscopic cholecystectomy or percutaneous catheter drainage, both to be performed within 24 hours after randomisation. A central study coordinator carried out randomisation using an online module and permuted block randomisation with varying block sizes with a maximum block size of four patients. Randomisation was stratified according to treatment centre. Owing to

the invasive nature of the intervention and the logistics involved in carrying out the procedures, neither the trial participants nor the investigators could be masked to group allocation.

### Procedures

Laparoscopic cholecystectomy was performed by the four trocar technique, with transection of the cystic duct and artery after reaching the critical view of safety, as described in national and international guidelines.<sup>13 14</sup> The procedures were performed by surgeons experienced in laparoscopic surgery, defined as performing more than 100 laparoscopic procedures yearly. Patients received a single dose of preoperative antibiotic prophylaxis according to the local hospital protocol.

Percutaneous catheter drainage was performed under local anaesthesia and aseptic circumstances, with image guidance using either ultrasonography or computed tomography. The procedures were performed by, or under direct supervision of, qualified radiologists. A certain experience to undertake this procedure was not required, as percutaneous catheter drainage is reported to be a relatively easy procedure, performed by any radiologist in the Netherlands. Gallbladder puncture was directed through the transhepatic or transperitoneal route, depending on the preference of the radiologist and the location of the gallbladder. For placement of the pigtail catheter into the gallbladder, either the trocar technique (one step technique) or the Seldinger technique (multiple step technique) was used. Emergency cholecystectomy was performed in case of clinical deterioration, persisting fever, or an increase in serum white blood cell count or C reactive protein within 48 hours, despite accurate position and function of the drain. Patients were discharged with the percutaneous drain. The drain was left in place for three weeks. Before removal of the drain, antegrade cholangiography was performed to assess for duodenal backflow and a patent cystic duct. Further treatment was left to the discretion of the treating clinician.

### Data collection and outcomes measures

The primary endpoints were death within one year and the occurrence of major complications, defined as infectious and cardiopulmonary complications within one month, the need for reintervention within one year, or recurrent biliary disease within one year. Table 1 provides detailed definitions. Reinterventions were either surgical, radiological, or endoscopic and had to be directly or indirectly related to acute cholecystitis. Routine elective cholecystectomy after percutaneous catheter drainage was not included in the primary endpoint (ie, only cholecystectomies for recurrent gallstone related complications were considered as matching the endpoint “need for reintervention”). Recurrent biliary disease was considered an endpoint only if readmission was required (so we did not include patients with recurrent biliary disease presenting to the general practitioner or emergency department without subsequent admission to hospital).

**Table 1 | Definitions of primary endpoints**

Endpoint	Definition	Comment
Death		Within one year after randomisation
Major complications:		
Intra-abdominal abscess	Fever or increased C reactive protein level/white blood cell count, or both, and intra-abdominal fluid collection on computed tomography or ultrasonography	Within 30 days after randomisation
Pneumonia	Coughing or dyspnoea, radiography with infiltrative abnormalities, increased infection variable, and positive sputum culture result	Within 30 days after randomisation
Myocardial infarction	Symptomatic increased cardiac enzyme levels and abnormalities on electrocardiography or cardiac ultrasonography	Within 30 days after randomisation
Pulmonary embolism	Radiologically proven pulmonary embolism	Within 30 days after randomisation
Need for reintervention	Surgical, endoscopic, or radiological reintervention	Within one year after randomisation. Before analysis, the adjudication committee decided to only report reinterventions directly or indirectly related to acute cholecystitis. A routine elective cholecystectomy after percutaneous catheter drainage was not included in the primary endpoint (ie, only cholecystectomies for recurrent gallstone related complications were included)
Recurrent biliary disease		Within one year after randomisation. Before analysis, the adjudication committee decided to only report recurrent biliary disease when readmission was required

The secondary endpoints included the individual components of the primary outcome, minor complications, difficulty of cholecystectomy (as scored by a visual analogue scale from 1 to 10), utilisation of healthcare resources, and total costs. The supplementary appendix provides details on cost calculation.

Follow-up took place at the outpatient clinic three weeks after discharge and subsequently by a phone call once every month for one year. Local clinicians performed data collection using case record forms. The study coordinator verified all completed forms in accordance with onsite source data. Through consensus, two investigators not involved in patient care resolved discrepancies detected by the study coordinator.

An adjudication committee consisting of four experienced surgeons and one radiologist carried out a blinded assessment of primary and secondary outcomes. Committee members individually evaluated the data in a standardised format for every patient, including all available data collected during follow-up. Disagreement was resolved in a plenary consensus meeting, with concealment of the treatment assignment.

### Statistical analysis

The sample size calculation was based on an expected reduction in the primary endpoints from 28% in the percutaneous drainage group to 15% in the cholecystectomy group.<sup>10</sup> This was based on the results of a Dutch retrospective cohort study and a systematic review analysing the safety and effectiveness of percutaneous drainage in elderly and critically ill patients.<sup>6-15</sup> To show this effect with 80% power, a two sided  $\alpha$  level of 5%, and a loss to follow-up of 1%, we needed at least 284 patients in total.

Primary analyses were performed in accordance with a pre-established analysis plan and according to the intention to treat principle. Differences between groups were expressed as risk ratios with corresponding 95% confidence interval. For continuous variables,

we calculated differences with the student's *t* test for normally distributed data and the Mann-Whitney U test for non-normally distributed data. We considered a two tailed P value <0.05 to be significant. All P values are two sided and not corrected for multiple testing.

In the original study protocol,<sup>11</sup> an interim analysis for efficacy was specified after the first year of inclusion because it was anticipated that half the number of required patients would have been randomised at that time. The accrual rate was, however, slower than expected. Therefore, we conducted the interim analysis at a later stage, when half the sample size had been reached. We compared the occurrence of the primary endpoints between the treatment groups. The Peto approach was followed, meaning that the study would only be stopped for beneficial effects in case of a P value <0.001. The independent data safety monitoring board evaluated the results of the interim analysis.

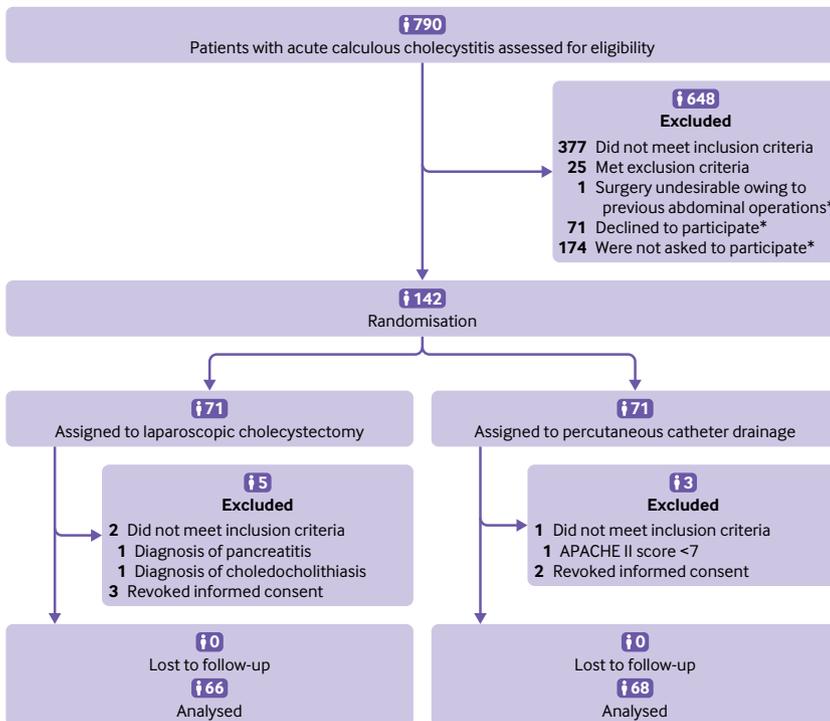
### Patient and public involvement

No patients were involved in setting the research question or developing plans for design of the study, nor were they asked to advise on interpretation or writing up of results. Results of the trial will be made available to all participants by email.

### Results

In December 2015, a formal interim analysis for the primary endpoints was performed. Data on 138 patients were reviewed, 118 of whom had completed follow-up. The P value of the difference between both groups was below the prespecified threshold of 0.001. Motivated by considerations of the beneficial effect of laparoscopic cholecystectomy and concerns about negative outcomes in the percutaneous drainage group, on 26 February 2016 the data safety monitoring board recommended termination of the trial. We followed up all patients who had undergone randomisation before this date until study completion.

Between 23 February 2011 and 30 January 2016 we assessed a total of 790 patients with acute calculous cholecystitis for eligibility, 142 of whom underwent



**Fig 1 | Enrolment, randomisation, and follow-up of study participants.** APACHE=acute physiology and chronic health evaluation. \*Patients who were eligible for inclusion but did not participate in the trial. The baseline characteristics of these patients were similar to those of the included patients, with the exception of the APACHE II score, which was higher in the included group (9.4 v 9.0) (see table 2 in supplementary appendix)

randomisation and completed follow-up (see results section and table 2 in the supplementary appendix). Eight patients were excluded from subsequent analyses because they revoked informed consent or did not meet the inclusion criteria in retrospect (fig 1). Baseline characteristics of the treatment groups were similar, with the exception of mean age (cholecystectomy 71.4 (SD 10.6) v 74.9 (SD 8.6)) and the number of patients with cardiovascular disease (58% v 78%; table 2).

Laparoscopic cholecystectomy was performed in 64 out of 66 participants assigned to this group. One patient underwent endoscopic retrograde cholangiopancreatography because of concomitant cholangitis, and one patient was treated conservatively because of hyponatraemia. Both underwent elective cholecystectomy several weeks after discharge. In 11 patients (17%), the laparoscopic procedure had to be converted to an open cholecystectomy. The median difficulty of the operation as scored by the performing surgeon was 8 (interquartile range 6-8). (See the results section in the supplementary appendix for details.)

Percutaneous catheter drainage was performed in all 68 patients assigned to this group. The procedure was technically successful in 65 patients (96%). In three patients the radiologist failed to place the percutaneous tube into the gallbladder lumen; two of these patients were treated conservatively until resolution of symptoms and one required emergency cholecystectomy owing to gallbladder perforation with extravasation of contrast fluid resulting in severe

abdominal pain. Clinical improvement within 48 hours occurred in 63 of the 68 patients (93%). In one patient an emergency cholecystectomy was performed as a result of clinical deterioration.

The rate of death did not significantly differ between the two groups; two patients (3%) in the cholecystectomy group and six (9%) in the drainage group died (P=0.27). Deaths in the cholecystectomy group occurred during follow-up and were related to oesophageal and colorectal cancer. In the drainage group two patients died during index admission as a result of ongoing sepsis due to the acute cholecystitis, one patient died during readmission from sepsis due to recurrent cholecystitis, and one patient died at home by an unknown cause, one week after removal of the percutaneous drain. The remaining two patients died during follow-up from mesothelioma and intestinal ischaemia.

Major complications occurred in eight of the 66 patients (12%) assigned to cholecystectomy and in 44 of the 68 patients (65%) assigned to drainage (risk ratio 0.19, 95% confidence interval 0.10 to 0.37; P<0.001) (table 3). These results did not change after post hoc adjustment for baseline differences in age and cardiovascular disease using multivariable logistic regression (adjusted odds ratio with cholecystectomy 0.08, 95% confidence interval 0.03 to 0.19; P<0.001). A formal test of interaction in a logistic regression model was used to assess whether treatment effects for the primary endpoints differed between subgroups based on treatment centre, and showed no significant difference (P>0.05).

The difference in rate of infectious and cardiopulmonary complications as well as minor complications between the two groups was not statistically significant (table 3). Reinterventions related to cholecystitis, however, were performed less often after cholecystectomy than after drainage (12% v 66%, P<0.001) (table 3). Recurrent biliary disease also occurred less often in patients assigned to cholecystectomy (5% v 53%, P<0.001) (table 3). Emergency cholecystectomy was performed in 11 of the 68 patients (16%) assigned to percutaneous catheter drainage—in two patients (3%) because of clinical deterioration and in nine (13%) because of recurrent cholecystitis (table 3). Elective cholecystectomy was performed in 20 of the 68 patients (29%) in the drainage group; in 15 patients (22%) because of recurrent gallstone related complications and in five (8%) because of either a dysfunctional drain or absence of duodenal backflow revealed by cholangiography. The indication for cholecystectomy in the latter five patients was, in the opinion of the adjudication committee, debatable, and therefore not considered as reaching the primary endpoint. Biliary injury occurred in four patients (6%) in the cholecystectomy group and in two (3%) in the drainage group, all of whom required endoscopic retrograde cholangiopancreatography. The total length of hospital stay (including readmissions) was five days (interquartile range 4-8 days) in the cholecystectomy group and nine (6-19) days in the

**Table 2 | Baseline characteristics of participants. Values are numbers (percentages) unless stated otherwise**

Characteristics	Laparoscopic cholecystectomy (n=66)	Percutaneous catheter drainage (n=68)
Mean (SD) age (years)	71.4 (10.6)	74.9 (8.6)
Men	41 (62)	44 (65)
Mean (SD) body mass index*	28.7 (5.3)	29.0 (5.5)
Coexisting conditions:		
Cardiovascular disease	38 (58)	53 (78)
Pulmonary disease	13 (20)	14 (21)
Chronic renal insufficiency	3 (5)	5 (7)
Diabetes	13 (20)	16 (24)
Previous abdominal surgery	16 (24)	10 (15)
ERCP before randomisation	3 (5)	4 (6)
ASA classification on admission:		
I: healthy status	10 (15)	4 (6)
II: mild systemic disease	33 (50)	37 (54)
III: severe systemic disease	23 (35)	24 (35)
IV: severe systemic disease that is a constant threat to life	0	3 (4)
Disease severity:		
Mean (SD) APACHE II score†	9.5 (1.9)	9.4 (2.0)
Mean (SD) C reactive protein level (mg/L)	218.5 (117.2)	214.7 (123.8)
Mean (SD) white blood cell count ( $\times 10^9/L$ )‡	17.0 (5.1)	17.2 (5.2)
Mean (SD) body temperature (°C)§	37.7 (1.1)	37.8 (0.9)
Median (interquartile range) time since onset of symptoms (days)¶	3 (2 to 3)	2 (1 to 4)

ERCP=endoscopic retrograde cholangiopancreatography; ASA=American Society of Anaesthesiologists; APACHE II=acute physiology and chronic health evaluation II.

\*Data missing for 12 patients in cholecystectomy group and nine in drainage group.

†Scores on acute physiological and chronic health evaluation II (APACHE II) scale range from 0 to 71, with higher scores indicating more severe disease.

‡Data missing for one patient in drainage group.

§Data missing for five patients in cholecystectomy group and two in drainage group.

¶In all patients, time since onset of symptoms was seven days or less. Data on exact number of days were missing for five patients in drainage group but were reported to be less than 7.

percutaneous drainage group ( $P<0.001$ ) (table 3). The total number of visits to the emergency department was seven and 56, respectively ( $P<0.001$ ). The total number of readmissions was nine in the cholecystectomy group and 67 in the drainage group ( $P<0.001$ ), and the total number of reinterventions was 21 and 64, respectively ( $P<0.001$ ).

The mean direct medical costs per patient during a follow-up of one year after randomisation were £4993 (\$6125; €5568) for cholecystectomy and £7427 for drainage, with a mean absolute difference of £2434 per patient. Details of costs are given in table 1 in the supplementary appendix.

## Discussion

This study showed that laparoscopic cholecystectomy is superior to percutaneous catheter drainage in the treatment of high risk patients with acute calculous cholecystitis. Cholecystectomy not only reduced the rate of major complications but also reduced utilisation of healthcare resources and costs by more than 30%.

Previous studies found a high short term success rate from percutaneous catheter drainage for acute cholecystitis in high risk patients.<sup>6 16-21</sup> Our study supports these findings, with prompt clinical improvement in more than 90% of patients undergoing percutaneous drainage. A systematic review of 53 mostly retrospective studies, published in 2009,<sup>6</sup> analysing the safety and efficacy of percutaneous catheter drainage in elderly and critically ill patients with acute cholecystitis, found few complications (6%)

and low procedure related mortality (0.4%), which corresponds with our finding. The reported 30 day mortality was high (13%), which could be attributed to confounding by indication, as most patients treated with drainage included in the retrospective studies are generally in poor clinical condition. In our study, the rate of death in the drainage group was also high (9%), but was not significantly different from that in the cholecystectomy group. The high rate of recurrent gallstone related disease in our study is reason for concern. More than half of the patients in the drainage group developed recurrent symptoms requiring (emergency) readmissions or reinterventions, or both; a much higher rate than reported in previous studies.<sup>17 18 22 23</sup> This may be explained by the randomised design of our study and the fact that we only included high risk patients.

The rate of recurrent gallstone related symptoms after drainage could have been lower if all patients would have undergone elective cholecystectomy. Routine elective cholecystectomy was not part of the study design because one of the advantages of percutaneous catheter drainage is the avoidance of complications related to surgery. For that reason, several authors suggest that high risk patients should not undergo elective cholecystectomy after percutaneous catheter drainage.<sup>20 24-26</sup> Our findings, however, support routine cholecystectomy in all patients who have undergone percutaneous catheter drainage, as promoted by others.<sup>27-29</sup> Alternatively, only patients who are especially at risk for recurrent gallstone related disease could undergo cholecystectomy. No studies, however,

**Table 3 | Primary and secondary endpoints for participants allocated to laparoscopic cholecystectomy or percutaneous catheter drainage. Values are numbers (percentages) unless stated otherwise**

Outcomes	Cholecystectomy group (n=66)	Drainage group (n=68)	Risk ratio (95% CI)	P value
<b>Primary endpoints*</b>				
Death	2 (3)	6 (9)	0.34 (0.07 to 1.64)	0.27
Major complication†	8 (12)	44 (65)	0.19 (0.10 to 0.37)	<0.001
<b>Secondary endpoints*</b>				
Death:	2 (3)	6 (9)	0.34 (0.07 to 1.64)	0.27
Directly/indirectly related to acute cholecystitis	0	3 (4)		
Unrelated to acute cholecystitis	2 (3)	2 (3)		
Unknown cause	0	1 (2)		
Infectious and cardiopulmonary complication‡:	5 (8)	3 (4)	0.97 (0.89 to 1.05)	0.49
Intra-abdominal abscess	4 (6)	2 (3)		
Pneumonia	2 (3)	1 (2)		
Myocardial infarction	0	0		
Pulmonary embolism	0	0		
Need for reintervention‡:	8 (12)	45 (66)	0.18 (0.09 to 0.36)	<0.001
Surgical intervention	3 (5)	32 (47)	0.10 (0.03 to 0.30)	<0.001
Emergency cholecystectomy	NA	11 (16)		
Clinical deterioration	NA	2 (3)		
Recurrent cholecystitis	NA	9 (13)		
Elective cholecystectomy§	2 (2)	20 (29)		
Recurrent gallstone related disease	NA	15 (22)		
Dysfunctional drain¶	NA	1 (2)		
Absence of duodenal backflow revealed by cholangiography¶	NA	4 (6)		
Cholecystectomy not performed during index admission¶	2 (2)	NA		
Diagnostic laparoscopy	1 (2)	1 (2)		
Endoscopic intervention	6 (9)	11 (16)	0.56 (0.22 to 1.43)	0.22
ERCP	6 (9)	11 (16)		
Cholelithiasis	2 (3)	9 (13)		
Biliary injury	4 (6)	2 (3)		
Removal of biliary stent	1 (2)	0		
Radiological intervention	4 (6)	15 (22)	0.28 (0.10 to 0.79)	0.008
Percutaneous catheter drainage	NA	8 (12)		
Recurrent cholecystitis	NA	6 (9)		
Dysfunctional drain	NA	2 (3)		
Drainage abscess	3 (5)	4 (6)		
Drainage biloma	2 (3)	0		
Drainage ascites	0	1 (2)		
Antegrade cholangiography	NA	4 (6)		
Contrast image PTC tube	1 (2)	0		
Recurrent biliary disease‡:	3 (5)	36 (53)	0.09 (0.03 to 0.27)	<0.001
Requiring emergency readmission	3 (5)	28 (41)		
Requiring planned readmission	2 (3)	16 (24)		
Elective cholecystectomy	NA	15 (22)		
ERCP	2 (3)	3 (4)		
Minor complication:	0	4 (6)		0.12
Wound infection	0	2 (3)		
Bleeding	0	0		
Urinary tract infection	0	2 (3)		
Healthcare utilisation:				
Median (interquartile range) length of stay after randomisation (days)	4 (3-6)	6 (4-8)		0.01
Median (interquartile range) total length of hospital stay (days)	5 (4-8)	9 (6-19)		<0.001
Median (interquartile range) total length of stay in ICU (days)	0 (0-0)	0 (0-0)		0.16
Total No (range per patient) of ER visits per study group	7 (0-1)	56 (0-5)		<0.001
Total No (range per patient) of reinterventions per study group	21 (0-6)	64 (0-4)		<0.001
Total No (range per patient) of readmissions per study group	9 (0-2)	67 (0-5)		<0.001

ERCP=endoscopic retrograde cholangiopancreatography; PTC=percutaneous transhepatic cholangiography; ICU=intensive care unit; ER=emergency department; NA=not applicable.  
\*Multiple events in same patient were considered as one endpoint.  
†For example, infectious and cardiopulmonary complications within one month, need for reintervention within one year, or recurrent biliary disease within one year.  
‡Included in primary endpoint of major complications.  
§Elective cholecystectomies were not included in primary endpoint, unless performed for recurrent gallstone related disease.  
¶These procedures were not included in primary endpoint because the adjudication committee judged the indication debatable.

have evaluated clinical, biochemical, or radiological predictors for failure of percutaneous catheter drainage in acute cholecystitis. Antegrade cholangiography may help to select patients who might benefit from elective surgery. Our study was not designed to evaluate the value of cholangiography after percutaneous catheter

drainage. Further prospective studies on this topic are needed.

Although other studies have reported considerable morbidity and mortality from emergency laparoscopic cholecystectomy in high risk patients (up to 41% and 5%, respectively),<sup>34 56</sup> we found that

immediate cholecystectomy in these patients is safe. Major complications occurred in 8% of patients, which seems acceptable in this category of severely ill patients. Yet it should be emphasised that the results of this trial only apply to patients with an APACHE II score of 7 or more and 14 or less, and so do not apply to patients with a score of 15 or more. During the study period, however, we only excluded 10 patients on the basis of this criterion. This implies that virtually all patients with acute calculous cholecystitis can safely undergo early laparoscopic cholecystectomy. As opposed to percutaneous catheter drainage, cholecystectomy is a definitive treatment for gallstone related disease, which does not require readmissions and other interventions that impact patient's quality of life and are a burden on hospital capacity for emergency and elective care. It may be clear that, in patients with a strict contraindication for surgery, percutaneous drainage is still an appropriate treatment, either as a bridge to surgery or as definite treatment.

### Conclusion

Among high risk patients with acute cholecystitis, laparoscopic cholecystectomy compared with percutaneous drainage is the preferred treatment strategy from both a clinical and economical point of view.

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The results of this trial were published in CSL's thesis entitled "Optimal treatment of acute cholecystitis." Utrecht University, Netherlands, 2017.

**Contributors:** PvD, MGHb, DJG, GAPN, JCK, SCD, AAWVG, PMK, DR, KK, AP, DvdP, RMPHC, BvR, TLB, and DB participated in the design of the study. CSL, PvD, MGHb, DJG, GAPN, JCK, SCD, AvG, PMK, DR, KK, VK, AP, DvdP, RMPHC, BvR, TLB, and DB collected data. CSL, HCvS, and JK did the statistical analysis. CSL, Hvc, JCK, and DB interpreted the data. CSL drafted the first version of the manuscript. CSL, HCvS, and DB drafted the subsequent versions of the manuscript. DB is guarantor. All authors read and appraised the report and approved the final version of the manuscript.

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**Ethical approval:** This study was approved by the institutional review board of the St Antonius Hospital, Nieuwegein, Netherlands (NL33662.100.10), as well as by the institutional review board of each participating centre. All patients gave informed consent.

**Data sharing:** A full dataset is available from the corresponding author at [d.boerma@antoniusziekenhuis.nl](mailto:d.boerma@antoniusziekenhuis.nl), contingent on approval from the institutional review board at the St Antonius Hospital, Nieuwegein, Netherlands. Consent was not obtained for data sharing but the presented data are anonymized and risk of identification is low.

**Transparency:** The lead author (DB) affirms that the manuscript is an honest, accurate and transparent account of the study being reported, and follows the CONSORT guidelines for the reporting of clinical trials. No important aspects have been omitted, and any discrepancies from the study as planned have been explained.

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**Supplementary information:** appendix