

Randomized controlled trial of laparoscopic cholecystectomy after ENGBD versus PTGBD in acute cholecystitis: the LCEVP study protocol

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Study protocol

Keywords: acute cholecystitis, endoscopic naso-gallbladder drainage, cholecystectomy

Posted Date: August 19th, 2019

DOI: <https://doi.org/10.21203/rs.2.13105/v1>

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Abstract

Background Transitional drainage which is followed by cholecystectomy plays a key role in the management of acute cholecystitis, especially in high-risk surgical patients. Endoscopic naso-gallbladder drainage (ENGBD) is an alternative to percutaneous transhepatic gallbladder drainage (PTGBD) for patients who need temporary drainage. There is a lack of prospective comparison on the relevant outcomes of the two drainage methods during the period of drainage, especially the subsequent cholecystectomy. **Methods** This is a randomised controlled two arm double-blind single center trial. Patients with acute cholecystitis undergo emergent or early cholecystectomy and need drainage will be randomly assigned to group PTGBD or ENGBD. Pain score is defined as the primary endpoint, whereas several secondary endpoints such as the rates of technical success, clinical remission, open conversion of cholecystectomy will be determined to elucidate more detailed differences between two groups. The general feasibility, safety and quality checks required for high quality evidence will be adhered to. **Discussion** This study would provide the first type A evidence concerning the comparison of ENGBD versus PTGBD in surgically high-risk patients with acute cholecystitis, it will be the first trial designed to determine the impact of two drainage methods on not only peri-drainage but also peri-LC. Trial registration NCT03701464. Registered on 10 October 2018.

Background

Laparoscopic cholecystectomy (LC) was the fundus-first approach for acute cholecystitis (AC). As the global population ages, not only the incidence of AC increases because of the strong relation between AC and age by previous studies,(1, 2) but an increasing number of patients with comorbidities who are poor surgical candidates require precise individualized management. According to Tokyo Guidelines 2018(TG18) flowchart,(3) percutaneous transhepatic gallbladder drainage(PTGBD) should be considered as the first-line alternative to surgical intervention in high-risk patients, especially in some Grade II (moderate) and most of Grade III (severe) AC by the TG18 severity grading.(4) Nevertheless, it may be limited in patients with thrombocytopenia, coagulopathy, large ascitic fluid, and Chilaiditi syndrome.(5, 6) In this circumstance, endoscopic gallbladder drainage (EGBD) had also been made another alternative management in high-volume institutes by skilled endoscopists by TG18 management strategies for gallbladder drainage.(7) EGBD, including endoscopic transpapillary gallbladder drainage (ETGBD) by using naso-gallbladder drainage (ENGBD) or gallbladder stent (EGBS) and endoscopic ultrasound-guided gallbladder drainage (EUS-GBD), is an effective method for acute cholecystitis both technically and clinically and seems to be safer than traditional PTGBD by a systematic review.(8)

Despite a prospective study comparing EUS-GBD and PTGBD and a randomized, controlled trial had confirmed that ENGBD and EGBS all appear to be suitable for the drainage(9)so far, there is no prospective comparison of ENGBD or EGBS with PTGBD, especially in their impacts while LC. Temporary or permanent drainage is essential for AC patients at high risk of surgery. In this study, we regard drainage as a temporary bridge to cholecystectomy in patients at high risk of early LC, yet stents were not expected to fall off during the waiting window, we chose this untouched territory comparing the ENGBD and PTGBD.

PTGBD has a technical success rate of nearly 97%, clinical response rates range from 56% to 100%,(6) which is also associated with an overall adverse events rate as high as 14%, including bile leak peritonitis, cholecystitis, bleeding, subcapsular hematoma, pneumothorax, catheter misplacement, and inadvertent removal. Moreover, it is contraindicated in patients with large peritoneal ascites, coagulopathy, thrombocytopenia and Chilaiditi syndrome. (5) For these reasons, less-invasive forms of ENGBD that are more efficient, more safe and less painful have been developed.(10) It was first proposed in 1984, but reported in 1990,(11) the technical success, clinical success, post-

procedure adverse events were 88.9%, 81.5%, 9.3% respectively according to the data provided by the only two randomized controlled studies(9, 12) of comparing ENGBD and EGBS. Despite such promising achievement from previous studies, the benefits of ENGBD and PTGBD on major clinical outcomes have never been compared prospectively. In particular, the effects of the two drainage method on LC remains uncertain and no extensive evaluation but only a retrospective study pointed out that ENGBD was more conducive to the smooth implementation of LC.(13)

Methods

Primary objective

The primary objective of the study is to determine the clinical pain remission of ENGBD versus PTGBD in patients requiring temporary gallbladder drainage, our hypothesis is that in comparison with standard PTGBD, ENGBD could not only relieve the pain symptoms associated with acute cholecystitis itself, but also relatively alleviate the severe pain caused by drainage.

Secondary objectives

The study will also compare ENGBD and PTGBD on clinical outcomes of drainage and the difficulties of cholecystectomy in surgically high-risk patients with acute cholecystitis, especially drainage success rate, clinical remission, adverse events. Also, their impacts on cholecystectomy will be focused and targeted, including difficulty grading of cholecystectomy, cholecystectomy duration, pathology and so on.

Trial design

The LCEVP is an investigator-initiated, parallel group, single-center, randomized controlled, with the allocation ration of 1:1, non-blind trial, however, the statistician would not know the grouping.

Participants, interventions, and outcomes

This manuscript was written adheres to SPIRIT 2013 Statement.(14) The checklist is available as an attached file1.

Study setting

The trial would be set at an university affiliated tertiary-care A hospital center which performs approximately more than 2000 ERCPs per year. The outline of trial procedures is presented in Figure 1.

Eligibility criteria

Inclusion criteria

Patients that were not considered suitable for early or urgent cholecystectomy because of high surgical risk, in which case biliary drainage followed by delayed LC is recommended by the TG18 flowchart: Grade II (moderate) AC, antibiotics and general supportive care fail to control inflammation; Grade III (severe) AC, after antibiotics and general organ support, negative predictive factors present including jaundice (TBil ≥ 2), neurological dysfunction, respiratory dysfunction or no negative predictive factors present but American Society of Anesthesiologists physical status classification (ASA-PS) is 3 or greater or Charlson comorbidity index (CCI) is 4 or greater.

Exclusion criteria

Exclusion of patients who meet one or more of the following criteria: age <18 or >90 years, pregnant or breastfeeding, severe obesity (body mass index ≥ 35 kg/m²), consent refusal, coagulation dysfunction (INR > 1.5) and low peripheral blood platelet count ($<50 \times 10^9 / L$), using anticoagulation or antiplatelet drugs; bile duct stones; prior surgery of Bismuth \square , Roux-en-Y and choledochojejunostomy, preoperative coexistent diseases: acute pancreatitis, GI tract hemorrhage or perforation, severe liver disease (such as decompensated liver cirrhosis, liver failure and so on), any malignant diseases.

Endoscopic surgeons criteria

All ERCP operations and LC are completed by an endoscopic surgeons team, with more than 10 years of ERCP and 20 years of LC experience, 400 cases of ERCP and 450 cases of LC per year at present.

Interventions

A systematically trained experienced attending physician assesses that once evidence of gallbladder drainage is available, all patients eligible for inclusion were informed fully of written informed consent, then would be randomly assigned to the interventional group (ENGBD) or to the reference group (PTGBD). Of course, risks and benefits associated with surgery would be routinely communicated and agreed in writing.

The experimental group patients are sedated by intravenous administration of sufentanil and propofol, then selective bile duct cannulation, a 0.025- or 0.035-inch guidewire is advanced into the cystic duct and subsequently into the gallbladder, withdraw the catheter, a 5Fr nasogallbladder catheter is inserted into the gallbladder along the guidewire. PTGBD is guided by ultrasound, an 18-gauge needle is inserted into the gallbladder, a 0.035-inch guidewire is coiled into the gallbladder and a 9Fr dilator expands the skin, then an 8Fr 20cm catheter is placed.

When ENGBD was technically unsuccessful or clinically ineffective, endoscopic nasobiliary drainage (ENBD), PTGBD and EUS-GBD were used as alternative procedures. The patients of inclusion would be given appropriate and quality treatment at anytime, in order to reduce waiting and improve adherence. Properly trained volunteers would accompany patients on key treatments, such as ERCP and PTGBD, extubation, LC, etc. This not only improves adherence because of the convenience to patients, but monitor adherence to a certain extent.

All patients of inclusion would be administered by the attending physician in accordance with recent guidelines for AC management, including sufficient infusion, maintenance of electrolyte balance, antibacterial agents, the monitoring of respiratory and hemodynamics, correction of acidosis and complications.

In particular, there is no strict recommendation for extubation time, ENGBD group would extubate when the bile in the nasogallbladder tube becomes clear, which is approximately 4 days after the drainage based on our experience. PTGBD group extubate after more than two weeks due to fistula formation. Also, no consensus has been reached about the optimal timing of LC after drainage. Because most studies(15–17) figure out the short interval increases intraoperative difficulty, we require all patients to undergo cholecystectomy 3 months after drainage, so that edema and inflammation around the gallbladder subsided completely.

Outcome

Primary outcome measures

The primary outcome is the pain score, defined as pain experienced during mobilization from the supine to the standing position. Using the visual-analogue scale, pain score would be obtained within 2 hours before drainage and at post-drainage 24h, 48h, 72h in conscious and communicating patients by a specially trained nurse, who devoted herself to the objectivity and authenticity of the scores. The details of assessment are as follows: draw a 10 cm line on a piece of paper, mark one end of the line with the number 0, indicating no pain; the other end with 10, indicating the most severe pain; the middle part indicates different degrees of pain. While assessing the pain of patients, make sure the patient could not see the numbers on the paper and let them mark the position according to their feelings about the pain. Then the nurse would get a score based on the mark. Pain scores reflect directly the degree of pain, the remission of pain not only relieves the pain symptoms associated with acute cholecystitis itself, but relatively alleviates the severe pain caused by tube.

In order to avoid the effects of anesthesia, we chose to evaluate the pain 24 hours after drainage. Especially, all pain assessment should be performed without the administration of pain killers or after using an analgesics for 6 hours. All modifications in pain management are recorded. Our team routinely prepares a multimodal approach to pain management, which is applied mainly based on nonsteroidal anti-inflammatory drugs (NSAIDs) and other adjuvant drugs, but tries to avoid applying it.

Secondary outcome measures

Table 1 presents details. These information comprehensively reflect effectiveness and safety of drainage, and difficulties in cholecystectomy.

Participant timeline

Please see table 2.

Sample size

The sample size calculation is based on the primary outcome-the pain score after drainage. According to the research by Jang, Ji Woong [et.al,\(18\)](#) using the method provided by Dehui Luo [et.al\(19\)](#) and Xiang Wan [et.al,\(20\)](#) the median post-procedure pain score in the PTGBD group was 5.4 ± 3.1 . And the mean visual analog score of post-

procedure pain in the ENGBD group was 1.3 by Takao Itoi [et.al.\(9\)](#) Using a theoretical sample size for two-sample design, 9 participants are needed in each group with 80% power and a 5% significance level. Allowing for a 10% loss to follow-up, we plan to recruit 22 patients (11 in each group).

Recruitment

The recruitment would last 20 months which began in June 2018. This duration was estimated based on the number of admissions for surgical high-risk AC patients at the team in previous 2 years. To achieve adequate participant enrollment, relevant work on the study would be performed around the clock, including nights and weekends as routine clinical practice.

Assignment of interventions

Allocation sequence generation and randomization

The randomization numbers were generated by statistician using computer program with 1:1 allocation, stored and encoded 20 sealed, opaque envelopes independently. When a participant meets inclusion and inclusion criteria, two volunteers first record the name and the code of the next unopened envelope, then open the envelope and sign their names. At last, the treatment assignment would be sent to the attending physicians, which could get only one allocation at a time after all pre-drainage preparations were completed, including the evaluation of patients and informed consent of participants.

Blinding

This is an open-label, unblinded trial for patients and physicians because of the nature of the intervention (position and shape of drainage tube). However, the investigator of collecting secondary outcomes, the assessor of clinical and biological data in charge of statistical analyses and outcome assessment will be masked as to the subjects assigned group.

Data collection, management, and analysis

Collection, management

All data are prospectively collected and managed by the well-trained research volunteers.

The baseline data would be collected and registered on inclusion: age, sex, BMI, body temperature, interval between onset of AC and drainage, comorbidities and coexisting conditions, ASA grade, CCI, AC grade based on TG 18 severity grading,(4) white blood cell (WBC) count, electrolytes, total bilirubin, CRP and so on. This would systematically reflect the characteristics of participants.

The peri-drainage data would be obtained from 2 hours before drainage to one week after ENGBD or PTGBD, such as drainage success, clinical remission, adverse events, pain scores within 2 hours before drainage and at post-

drainage 24h, 48h, 72h, etc.

The following data are collected and registered on peri-LC: gallbladder wall thickness changes (pre-drainage to pre-LC), duration and hemorrhage of LC, gallbladder integrity, drainage tube, hospitalization (hospital stay after gallbladder drainage and cholecystectomy), surgical site infection, severe adhesion, pathology, LC related complications.

Results that are negative or undetected or measured as 0 should have corresponding symbolic representation and can not be vacant, in order to distinguish from missing values. The treatment of outliers should be judged from both medical and statistical aspects by the statistician during blind examination.

Images and biological samples including venous blood and bile will be collected for probably ancillary and long-term research. Also, all patients will be followed up 6 months after cholecystectomy.

The volunteers of data acquisition have rich clinical experience and rigorous working attitude, and received training and guidance. Data storage would be carried out by two independent investigators to ensure reliability and validity.

We would also pay attention to and collect the information of the participants who discontinue or deviate before cholecystectomy and the follow-up are completed. To promote participants' retention and enthusiasm, we would establish workflow manuals contact patients regularly, communicate properly and take effective care of their health.

Analysis

A predefined statistical analysis plan will be followed. All statistical analyses were conducted with IBM SPSS (version 21.0). P values of less than 0.05 were considered to represent statistical significance. Categorical variables would be reported as counts and percentages. The mean and standard deviation or median with interquartile ranges (IQR) were used to describe continuous variable data, as appropriate.

The primary analysis is a comparison between ENGBD and PTGBD for the pain remission based on an intention-to-treat principle. A multilevel, random-slope model would be fitted to the data with time points nested in patients to allow for clustering of data within each patient. This model adjusted for the fixed effects of treatment group, time (-2, 24, 48 or 72 hours), and treatment × time interaction. An unstructured covariance pattern was selected for the repeated measurements as the least restrictive structure, which resulted in better model fit based on log-likelihood values than more constrained patterns. Estimates of the difference in pain scores between treatment groups were assessed overall and at individual time points.

The secondary analysis is a comparison between ENGBD and PTGBD for the rates of technical and clinical success, adverse events, gallbladder wall thickness changes. Also, the outcomes during cholecystectomy including duration time, open conversion would also be analyzed. The Chi-square test or Fisher exact test was used for dichotomous variables, and Wilcoxon test was used for rank data. For quantitative data, two-tailed Student's t test is used if the normal distribution and homogeneous total variance are satisfying, otherwise, Wilcoxon test is used.

A blinded adjudication committee will assess the occurrence of the primary and secondary endpoints after the last patient has completed follow-up.

Data monitoring

Setting up a data monitoring committee and interim safety analysis are unnecessary because of the high security and short cycle of this trial.

Harms

All adverse events thought to be related to the trial would be recorded rigorously and carefully. Any unexpected major serious complications suspected to be associated with ENGBD must be reported to interviewer and attending physicians, the trial may be temporarily stopped, and effective treatment measures in the first place will be taken by attending physicians.

Auditing

A auditing committee consisting of an endoscopist, a surgeon and a sonographer, would monitor the trial at least once every 3 months. They remain independent from investigators and audit the frequency, procedures, and safety of the trial.

Protocol amendments

Any changes to eligibility criteria, outcomes and analyses would obtain approval from the ethics committee and the clinical trial.

Confidentiality

Data would be handled in a confidential way. During the whole experiment, sensitive information such as the patient's name, identity card number would not be exposed by coding

Access to data

All investigators would have access to the final trial data set after the end of the study.

Dissemination policy

Data analysis, interpretation and findings would be presented at academic conference and published in peer-reviewed journals. After the end of the trial, the interviewer will complete the writing and submit the manuscript via open access as soon as possible. Every participant would get the results and conclusions by mail.

Discussion

To our knowledge, LCEVP would be the first randomised controlled study powered to investigate the effect of ENGBD and PTGBD on peri-drainage and peri-LC in poor surgical AC population. Moderate and severe AC are

usually associated with increased morbidity and mortality if rushed cholecystectomy is performed, it is becoming increasingly difficult to ignore the optimal management of AC, which includes transitional and permanent drainage, especially when it comes to critically ill patients with multiple comorbidities.

Only one small retrospective discovery of limited quality compared the ENGBD and PTGBD in AC patients to date. (13) Part of the reason may be that ENGBD has to precisely over-select into the difficult visualizing, long, narrow, tortuous cystic ducts, which is a challenge because the ducts could be blocked by stones. The weighted pooled rates for technical success of ENGBD is 81% according to a proportion meta-analysis.(8) However, this technique uses standard off-the-shelf ERCP accessories and can be performed in patients who are coagulopathic or using antiplatelet or anticoagulant agents, and it has certain advantages including better drainage effect which can be observed directly, convenient removal of drainage tube, no establishment of abnormal anatomic fistula and so on. Thus it may be regarded as a bridging therapy for acute cholecystitis followed by interval surgical cholecystectomy. Unfortunately, the impact of ENGBD on LC is rarely reported.

Compared to direct surgery, PTGBD shorten the operative duration of LC in patients with moderate to severe AC. (21) However it has a high risk of conversion to an open procedure during cholecystectomy by a multicenter analysis.(22)

As a large surgical endoscopy center that completed more than 2000 ERCPs per year, our team not only has rich experience in cholecystectomy, but also masters excellently the endoscopic minimally invasive technique, we are confident to complete this major impact study. The trial has the potential to optimize drainage management with direct and realistic benefits for critical AC patients. It is worth mentioning that we would also pay attention to the effects of the two drainage methods on the cholecystectomy. If the study finds out that ENGBD may contribute to late cholecystectomy, it will be an excellent alternative or a powerful complement to PTGBD.

Trial status

The current LCEVP protocol is version 3.0 (15 May 2019). The trial opened to recruitment on 10 October 2018. Maybe we would complete the recruitment on 10 October 2020.

List Of Abbreviations

ENGBD: Endoscopic naso-gallbladder drainage; PTGBD: percutaneous transhepatic gallbladder drainage; LC: Laparoscopic cholecystectomy; EGBD: endoscopic gallbladder drainage; ETGBD: endoscopic transpapillary gallbladder drainage; EGBS: endoscopic gallbladder stent; EUS-GBD: endoscopic ultrasound-guided gallbladder drainage; NSAIDs: nonsteroidal anti-inflammatory drugs.

Declarations

Ethics approval and consent to participate

The study is conducted in accordance with the Declaration of Helsinki, and registered at <http://www.clinicaltrials.gov> on 8 October 2018 with trial identification number NCT03701464. Ethics Committee of The First Hospital of Lanzhou University approved the study in 2018 and the approval Number is LDYYMENG2018–1027.

Access to consent and assent would be subject to as required by the Institutional Review Board in accordance with the 2013 Declaration of Helsinki. We would do everything possible to obtain the written consent of the patient who is informed about the trial. We also seek written informed consent from next of kin if the patient can not provide it because of severe illness, and continue the trial only when the patient agrees in writing consent after his/her condition is restored.

Consent for publication

Not applicable.

Availability of data and material

Not applicable.

Competing interests

The authors declare that they have no competing interests.

Sponsors and Funding

The lead sponsor is Hepatopancreatobiliary Surgery Institute of Gansu Province. This study was financially supported by the National Natural Science Foundation of China (Grant No. 81872036). The study sponsor and funders have had no input into study design; collection, management, analysis and interpretation of data; writing of the report; or the decision to submit the report for publication (beyond usual governance activities) and will not have any such input in the future.

Authors' contributions

All authors approved the final manuscript. PLM: protocol development, drafting of this manuscript, critical revision of the manuscript for significant intellectual content. PY, BB, YYL: providing personnel, environmental and financial support and tools and instruments that are vital for the project. HPW, JY: taking responsibility in logical interpretation and presentation of the results. TYL, YL: reviewing the article before submission not only for spelling and grammar but also for its intellectual content. WBM, XL: constructing an idea or hypothesis for manuscript, planning methodology to reach the conclusion.

Acknowledgements

The authors would like to thank Dr Hao Xu for giving us assistance during the therapeutic process.

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Tables

Table 1 Secondary outcome measures

Secondary outcome	Definition
Drainage success	Bile juice outflow more than 50ml from tube a day
Clinical remission	Improvement of typical clinical symptoms with laboratory tests, radiologic findings after the procedure
Adverse events	Migration, hemorrhage, perforation, bile leak in peri-drainage
Gallbladder wall thickness changes	The change of thickness of gallbladder wall pre-drainage and pre-lc
Severe adhesion	Establishing the critical view of safety is difficult
Duration	Time of laparoscopic cholecystectomy
Hemorrhage	The amount of bleeding during LC
Gallbladder integrity	Complete separation of the gallbladder from the liver, with no residual or damage
Drainage tube	The surgeon decides the placement of the abdominal tube
Surgical site infection	Superficial or deep incisional infections, or organ/space infections (i.e. intraabdominal abscesses)
Open conversion	Conversion to open surgery was decided by surgeon.
Pathology	Pathologic findings of excised gallbladder: necrotizing, suppurative, edematous, chronic cholecystitis.
LC-related complications	Bleeding, bile leakage, subhepatic abscess formation in peri-LC

Table 2 Participant timeline

STUDY PERIOD

TIMEPOINT	Enrollment	Allocation	Post-allocation											Close-out
	<12h	0	2h	12h	day1	day2	day3	day4	2w	2m	3m	4m	5m	6m
Eligibility screen	X													
Informed consent	X													
Laboratory test	X													
Allocation		X												
INTERVENTIONS														
ENGBD+LC				X	X	X	X	X			X			
PTGBD+LC				X	X	X	X	X	X		X			
ASSESSMENTS														
List baseline variables	X	X	X											
List pain score			X		X	X	X							
List second outcomes														
Peri-drainage						X	X	X	X					
Peri-LC										X	X			
Follow-up												X	X	X

Figures

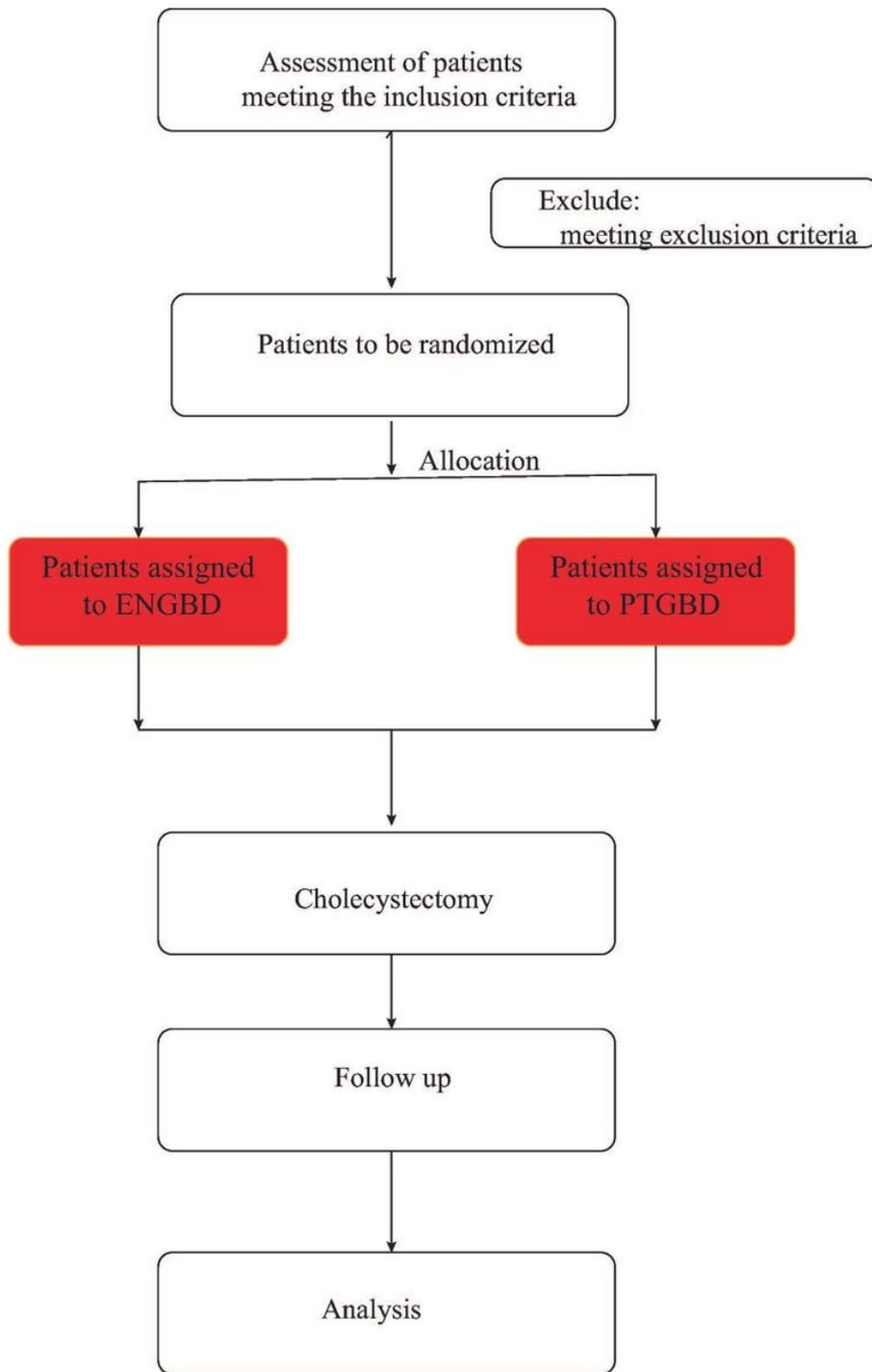


Figure 1

Flow chart of participants.

Supplementary Files

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