Laparoscopic and endoscopic cooperative surgery plus sentinel lymph node navigation surgery for elderly patients with early gastric cancer – phase II trial: Study protocol

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Abstract

Background: When patients are outside the criteria for endoscopic submucosal dissection (ESD) treatment, additional gastrectomy with radical lymph node dissection is generally performed. It is thought, however, that post-gastrectomy syndrome and decline in quality of life occur in elderly patients. In patients with submucosal-invasive gastric cancer who undergo gastrectomy, lymph node metastases (LNM) are very common. LNM in patients with submucosal-invasive gastric cancer after ESD, however, are very rare. In elderly patients who need additional treatment after ESD, partial gastrectomy with regional lymphadenectomy may be feasible in oncological. It is extremely difficult to determine the appropriate incision line in the laparoscopic view because of scarring in the intraluminal stomach after gastric ESD. We plan to perform laparoscopic and endoscopic cooperative surgery (LECS) to address this problem.

Methods: When performing LECS for patients with early gastric cancer (EGC), sufficient lymph node dissection is required, so we apply sentinel lymph node navigation surgery (SNNS). The purpose of this trial is to test the hypothesis that limited operation using LECS plus SNNS is effective and oncologically safe in the selected group including elderly patients who undergo ESD.

Conclusion: This study is a prospective, single arm, single center trial. Patients over 75 years old with EGC who are pathologically diagnosed outside the criteria for ESD treatment are eligible. The primary endpoint is to determine the rate of the postoperative complications. Secondary endpoints include determination of surgical results, postoperative outcomes, and survival rates.

Abbreviations: ESD: endoscopic submucosal dissection; LNM: lymph node metastasis; LECS: laparoscopic and endoscopic cooperative surgery; EGC: early gastric cancer; SNNS: sentinel lymph node navigation surgery; QOL: quality of life; ICG - indocyanine green; OS: overall survival; DFS: disease-free survival; SNs: sentinel lymph nodes; RCT: randomized controlled trial

Introduction

The number of patients with early gastric cancer (EGC) who undergo endoscopic treatment is increasing worldwide. Accurate diagnosis of invasive depth (clinical T factor) of gastric cancer is difficult [1], so in clinical practice, diagnostic endoscopic submucosal dissection (ESD) is often performed. If patients are outside criteria for ESD treatment, additional gastrectomy with radical lymph node dissection is generally performed [2]. It is generally accepted, however, that patients who undergo radical gastrectomy experience post-gastrectomy syndrome, such as appetite loss, epigastric discomfort, reflux symptom, weight loss, and decline in quality of life (QOL). Such post-gastrectomy syndromes appear remarkably in elderly patients [3,4].

After gastrectomy with submucosal-invasive gastric cancer, lymph node metastases (LNM) are very common (10 to 23%) [5-7]. LNM in patients after ESD are reportedly rare (0 to 5%), and good prognosis in patients with EGC treated with ESD may be obtained [8-10]. The site of LNM was the perigastric lymph nodes in all patients who underwent ESD [10]. Therefore, in elderly patients who need additional treatment after ESD, partial gastrectomy with regional lymphadenectomy may be oncologically feasible.

Scarring is present after gastric ESD in intraluminal stomach, so determining the appropriate incision line is extremely difficult by laparoscopic view. In recent years, laparoscopic and endoscopic cooperative surgery (LECS) was developed for gastric submucosal tumor [11] and the usefulness and feasibility were reported [12,13]. We therefore apply LECS to partial gastrectomy in patients with EGC after ESD and by minimizing the range of resection can improve postoperative QOL [14].

When performing LECS for patients with EGC, sufficient lymph node dissection is required to prevent recurrence of gastric cancer. If limited operation is adapted for EGC patients, increase in recurrence rate is not permissible, so lymph node dissection should be minimized by sentinel lymph node navigation surgery (SNNS). False negatives are a problem in conventional pigmentary visualization method [15]. Recently, however, there has been success in reducing false negative rate.
by using new modalities and tracers [16-18] particularly the method combining indocyanine green (ICG) and infrared light observation, which reduced the false negative rate to 0% [19,20]. The purpose of this trial is to test the hypothesis that limited operation using LECS plus SNNS (LECS-SNNS trial) is effective and oncologically safe in the selected group including elderly patients who undergo ESD.

LECS-SNNS trial is conducted in accordance with the Declaration of Helsinki. Wakayama Medical University Hospital Institutional Review Board approved the final version of the protocol (version 1.0) prior to the start of the study (approval number: 2235). The University Hospital Medical Information Network Clinical Trials Registry number is UMIN000031003 (https://upload.umin.ac.jp/cgi-open-bin/ctr_e/ctr_view.cgi?recptno=R000035311)

Materials and methods

**Objectives:** This trial aims to test the hypothesis that limited operation using LECS plus SNNS is effective and oncologically safe in the selected group, specifically elderly patients with EGC who were diagnosed as being outside the criteria for ESD treatment pathologically.

**Study setting:** The LECS-SNNS trial is a prospective, single arm, single center trial.

**End points:** The primary endpoint is to ascertain the rate of postoperative complications, which are analyzed by using the Dindo-Clavien classification [21]. Complications higher than grade II are regarded as clinically significant. Secondary endpoints are: (i) surgical results, such as operation time, blood loss, and transition rate to open surgery; (ii) postoperative course, such as time to start of drinking, time to start of eating, postoperative hospital stay, weight loss rate (version 1.0) prior to the start of the study (approval number: 2235). The University Hospital Medical Information Network Clinical Trials Registry number is UMIN000031003 (https://upload.umin.ac.jp/cgi-open-bin/ctr_e/ctr_view.cgi?recptno=R000035311)

**Inclusion criteria:** Patients over 75 years old with EGC selected to undergo additional resection after ESD are eligible for inclusion. Written informed consent is required.

**Exclusion criteria:**
1. History of preoperative chemotherapy for gastric cancer.
2. Ischemic heart disease requiring treatment.
3. Liver cirrhosis or active hepatitis
4. Severe respiratory disease (including interstitial pneumonia, pulmonary fibrosis, pulmonary emphysema).
5. Chronic renal failure requiring hemodialysis.
6. Other malignant disease that can influence adverse events.
7. Patients in whom laparoscopic surgery cannot be performed because of history of laparotomy in the upper abdomen.

**Registration:** After confirming fulfillment of the eligibility criteria, all handling cases are managed by subject identification code or anonymized registration number. The correspondence table of the anonymizing code and the name and the consent form containing the name are strictly kept in the separate lockable document storage of the Second Department of Surgery.

**Treatment methods**

**SNNS:** Under general epidural anesthesia, the patient is placed in the supine position with open legs and pneumoperitoneum of 10 mmHg is created. A laparoscope is inserted through the umbilical incision, and four trocars are placed in the upper abdomen. The overtube is inserted under the oral endoscope. We confirm the scar of endoscopic treatment and then a 0.5% ICG solution (Diagnogreen 0.5%; Daiichi Pharmaceutical, Tokyo, Japan) is endoscopically injected into the submucosa of the gastric wall at four sites (0.5 mL each) around the scar intraoperatively. After 20 minutes, ICG fluorescence imaging is carried out using the 1588 AIm camera system with Endoscopic Near - Infrared Visualization (Stryker Corporation, Kalamazoo, MI). Fluoresced lymph nodes are considered to be sentinel lymph nodes (SNs). SNs and surrounding sentinel lymphatic basin are extracted from the fatty tissue. Lymph nodes in the dissected specimens are isolated from the surrounding tissues on the back table. These lymph nodes are submitted for the intraoperative rapid medical examination (Figure 1).

**LECS:** Details of the LECS procedures performed at our institute have been previously described [12]. Blood vessels and omentum in the excision area around the scar are prepared using an ultrasonically activated scalpel (HARMONIC; Ethicon Endo-Surgery, Inc., Cincinnati, OH). The jejunum near the ligament of Treitz is clamped using detachable forceps to avoid air inflation in the gastrointestinal tract. With a view of the endoscopic image, a circumferential incision is made around the scar using an ESD technique involving a Flush Knife (Fujinon Optical Co., Ltd., Tokyo, Japan), which is used to perforate the stomach. The IT knife (Olympus, Tokyo, Japan) tip is inserted into the perforation site and seromuscular dissection is initiated along the incision line of the submucosal layer. With a view of the laparoscopic image, the seromuscular layer of the stomach is dissected along the incision line using HARMONIC, and the scar is removed. The post-excisional hole in the stomach is closed using a laparoscopic suturing technique.

If intraoperative pathologic diagnosis is metastatic lymph nodes, radical gastrectomy with D2 lymphadenectomy is performed (Figure 1).

**Postoperative management:** Nasogastric tube will be removed when awoken from anesthesia. Postoperative pain control consists of patient-controlled analgesia. Patients are encouraged to be out of bed and walking around the ward under the guidance of a physiotherapist or nurse the day after surgery. Patients will be discharged when they pass stool, are able to drink, can walk and are comfortable with oral analgesia.

**Follow-up:** All registered patients will be followed up for at least 5 years. Blood test, upper gastrointestinal endoscope, and enhanced abdominal computed tomography will be evaluated at least 5 years. Assessment of QOL will be performed preoperatively, at discharge, after six months, one year, two years, three years, four years, and five years postoperatively (Table 1).

**Study design and statistical analysis:** The planned sample size of the phase II study is 20 patients. Primary endpoint of LECS-SNNS trial is to determine the rate of postoperative complications. In the main analysis, the point estimate and the 95% confidence interval are constructed for the rate of postoperative complications. Total postoperative complications in radical gastrectomy in our department so far are 10% [4,25]. If the lower limit of the 95% confidence
interval exceeds 10%, LECS and SNNS is judged to be an unacceptable technique. Data will be collected prospectively for all patients including history, physical examination, laboratory data, pathologic examination, perioperative clinical information and complications. Data is collected via datasheets on paper and kept securely. All required parameters will be collected in a SPSS data file (SPSS version 25, IBM statistics Chicago, IL).

Discussion

We planned LECS-SNNS trial for patients over 75 years old with EGC selected to undergo additional resection after ESD. This LECS-SNNS trial is the first report of limited operation for elderly patients with EGC. If favorable outcomes are obtained in this study, this limited operation of LECS-SNNS may become a treatment option for elderly patients with EGC.

This trial has several limitations. It is a phase II study without randomized controlled trial (RCT) and is conducted at a single institution. Due to the small sample size, findings from this trial do not allow established clinical application, but rather serve to inform the need for larger multicenter phase III RCT.

Competing interests

No authors have competing interests to report.

Authors’ contributions

Maruoka designed the protocol and wrote the manual. Ojima further aided in assessment and revision of the protocol and revised the manuscript. All authors approved the final version of the protocol.

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