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**Laparoscopic spleen-preserving splenic hilar lymphadenectomy in consecutive 108 patients with upper gastric cancer**

Li P *et al*.Splenic hilar lymphadenectomy and gastric cancer

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**Abstract**

**AIM:** To evaluatethe feasibility and short-term efficacy of laparoscopic spleen-preserving splenic hilar (No. 10) lymphadenectomy in treating advanced upper gastric cancer (AUGC).

**METHODS****:** Between January and December 2012, 108 laparoscopic spleen-preserving No. 10 lymphadenectomy along with total gastrectomy with routine D2 lymphadenectomy were consecutively performed at our hospital to treat clinical T2-3 (cT2-3) upper gastric cancers. The preoperative clinical T stage was cT2 in 36 patients and cT3 in 72 patients. A prospectively designed database tracked the 108 patients, including the completeness of medical records and the adequacy of follow-ups. Patient clinicopathological characteristics, intraoperative and postoperative surgical outcomes, morbidity and mortality, lymph node (LN) dissection, and postoperative follow-up were analysed retrospectively.

**RESULTS:** Laparoscopic spleen-preserving No. 10 lymphadenectomy was successful in all 108 patients. The mean operation time was 169.3 ± 27.1 min, and the mean No. 10 lymphadenectomy time was 20.0 ± 5.7 min. The mean total blood loss was 46.2 ± 11.3 mL, and the mean blood loss from No. 10 lymphadenectomy was 14.3 ± 3.8 mL. The mean postoperative hospital stay was 11.9 ± 6.0 d. The intraoperative and postoperative morbidity rates were 3.7% and 12.0%, respectively, but there was no postoperative mortality. A mean of 44.4 ± 17.6 LNs were retrieved from each specimen, including 3.0 ± 2.4 No. 10 LNs. Three patients (2.8%) with cT3 cancer had LN metastasis of the splenic hilus, including two patients with pathological T3 (pT3) and one patient with pathological T4a (pT4a) tumours, all located in the greater curvature. No splenic hilar LNs metastasis was evident in the patients with pT1 and pT2 tumours. At the median follow-up time of 18 mo (range, 12 to 23 mo), all patients had survived and none had experienced recurrent or metastatic disease.

**CONCLUSION:** Laparoscopic spleen-preserving No. 10 lymphadenectomy is feasible and effective for treating AUGC. Routine No. 10 lymphadenectomy may be unnecessary for AUGC without serosa invasion unless T3 tumours are located in the greater curvature.

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**Key words:** Stomach neoplasms; Spleen-preservation; Laparoscopy; Gastrectomy; Lymphadenectomy

**Core tip:** Several studies have shown that laparoscopic spleen-preserving No. 10 lymphadenectomy is feasible for patients with upper gastric cancer, but the sample sizes in these studies were small. Furthermore, the value of the procedure must be further evaluated with large sample studies, and it is debatable whether routine No. 10 lymphadenectomy should be performed for advanced upper gastric cancer (AUGC) without serosa invasion. We have, therefore, evaluated the feasibility and short-term efficacy of laparoscopic spleen-preserving No. 10 lymphadenectomy in consecutive 108 patients with AUGC (cT2-3). In addition, early follow-up results will also be presented.

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**INTRODUCTION**

Splenic hilar lymph nodes (No. 10 LNs) are the LNs located in the splenic hilum, including those LNs adjacent to the splenic artery distal to the pancreatic tail, on the roots of the short gastric arteries, and along the left gastroepiploic artery proximal to the first gastric branch[1]. Standard D2 LN dissection during total gastrectomy for advanced upper gastric cancer (AUGC) requires the removal of the No. 10 LNs[1]. Spleen-preserving No. 10 lymphadenectomy has been shown to be technically feasible and safe for patients undergoing open surgery for upper gastric cancer[2], with lower postoperative morbidity and mortality rates than splenectomy[3-7]. Moreover, the radical effects and long-term survival rates were similar to those effects and rates in patients who underwent splenectomy[8-12]. Total gastrectomy with spleen-preserving No. 10 lymphadenectomy is, therefore, increasingly used to treat patients with upper gastric cancer. Spleen-preserving D2 lymphadenectomy, however, requires an anatomical No. 10 lymphadenectomy, a procedure that is technically difficult because of the presence of intricate and complex vessels and a narrow and deep space at the splenic hilum. Moreover, complete removal of No. 10 LNs is particularly difficult in obese patients and patients with splenic adhesions.

Laparoscopic gastrectomy with D2 lymphadenectomy has been shown to be safe and feasible for patients with advanced gastric cancer[13-16]. Although several studies have shown that laparoscopic spleen-preserving No. 10 lymphadenectomy is feasible during total gastrectomy with D2 LN dissection for small numbers of patients with upper gastric cancer [17-19], few large-scale studies have assessed its success in patients with AUGC. Furthermore, it is debatable whether routine No. 10 lymphadenectomy should be performed during total gastrectomy for AUGC without serosa invasion. In the current study, we have, therefore, evaluated the feasibility and short-term efficacy of laparoscopic spleen-preserving No. 10 lymphadenectomy in consecutive 108 patients with cT2-3 AUGC. In addition, early follow-up results will also be presented.

**MATERIALS AND METHODS**

***Patients***

Between January and December 2012, consecutive 108 patients with cT2-3 AUGC underwent laparoscopic spleen-preserving No. 10 lymphadenectomy, along with total gastrectomy and routine D2 lymphadenectomy in the Department of Gastric Surgery, Fujian Medical University Union Hospital. Beginning in May 2007, the surgeon (Huang Chang-Ming) in this study had performed more than 500 laparoscopy-assisted gastrectomies with D2 LN dissection in gastric cancer patients before attempting this procedure. Since then, a prospectively designed database has tracked all laparoscopy-assisted gastrectomies for gastric cancer. Moreover, the surgeon performed laparoscopic spleen-preserving No. 10 lymphadenectomy for advanced proximal gastric cancer using a left-sided approach[20]. This method was mastered after a learning curve of 40 patients[21]. The group of consecutive 108 patients was used for our retrospective analysis because of the completeness of their medical records and the adequacy of their follow-ups. Upper gastric cancer was diagnosed by an analysis of endoscopic biopsy specimens. Preoperative imaging studies were routinely performed following endoscopic examination, computed tomography (CT) scanning, abdominal ultrasonography (US) and endoscopic US. CT scans and multi-slice spiral CT angiography (MSCTA) were performed to preoperatively assess the splenic vascular anatomy(Figures 1 and 2). Advanced (cT2-T3) upper gastric cancer diagnosed by preoperative CT scanning and endoscopic US were enrolled in this study. Patients with clinical T1 (cT1) or clinical T4 (cT4) tumors, distant metastasis, or preoperative enlargement or integration of LNs were excluded. No patients had received preoperative chemoradiation therapy. Each preoperative patient was informed of the surgical procedure, including its advantages and risks. All patients provided written informed consent for the procedure prior to surgery as well as for the publication of this report and any accompanying images.

In the current study, the No. 10 lymphadenectomy began with the surgeon using an ultrasonic scalpel to separate and reveal the end of the splenic arteries within the retropancreatic space at the superior boder of the pancreatic tail, until to divide the last short gastric artery. The No. 10 lymphadenectomy time referred to the time of this procedure. The blood loss during operation was measured by estimating the volume of blood in the suction container and weighing the gauze with blood. Dissected LNs were classified according to the 3rd English edition of the Japanese classification of gastric carcinoma[1]. The clinical and pathological stagings were in accordance with the American Joint Committee on Cancer (AJCC) seventh edition of Gastric Cancer TNM Staging[22]. Follow-up was performed by trained investigators every 3 mo. The patients’ routine follow-ups consisted of a physical examination, laboratory tests, chest radiography, abdominopelvic ultrasonography or CT scans. The survival time was calculated from the time of surgical intervention until the last date of contact (December 31, 2013).

***Surgical procedures***

**Patient positioning**: The patient was placed in the reverse Trendelenburg position with the head elevated approximately 15° to 20°, and tilted with the left side up approximately 20° to 30°. A 10-mm trocar for the laparoscope was inserted below the umbilicus; a 12-mm trocar was inserted in the left upper quadrant as a major hand port; a 5-mm trocar was inserted in the left lower quadrant as an accessory port; a second 5-mm trocar for exposure was inserted in the left upper quadrant, and a third 5-mm trocar for exposure was inserted in the right lower quadrant. The surgeon stood on the left side of the patient; the assistant surgeon was on the right side, and the camera operator was situated between the patient’s legs.

**Other lymphadenectomy:** The gastrocolic ligament was divided using an ultrasonic scalpel along the border of the transverse colon. The right gastroepiploic vein and the right gastroepiploic artery were vascularised and divided to dissect the No. 6 LNs. The stomach was lifted toward the head to expose the gastropancreatic fold. The LNs along the proximal splenic artery (No. 11p) at the upper border of pancreatic body were removed. Next, the dissection continued rightward. The fatty connective tissue, including the LNs along the celiac trunk (No. 9), the left gastric artery (No. 7), and the common hepatic artery (No. 8a) were removed en-block with the left gastric vein and the left gastric artery being vascularised and divided. Then, the LNs around the right gastric artery (No. 5) and along the surface of the proper hepatic artery (No. 12a) were dissected and removed. Subsequently, the liver was held up to divide hepatogastricum ligament along the lower border of the liver and the LNs around lesser curvature (No. 3) was removed. At last, the phrenoesophageal membrane and both vagus nerves were divided and the LNs around the abdominal esophagus (No. 1 and 2) were dissected.

**No. 10 lymphadenectomy:** The patient was subsequently tilted with the left side up approximately 20° to 30° and subjected to a 20° upward head tilt. The surgeon then moved to stand between the patient’s legs, and the assistant and camera operator were both on the patient’s right side. Before surgery, the assistant placed the greater omentum behind the stomach to keep the visual field clear, pulled and tensed the gastrosplenic ligament. The surgeon gently pressed the tail of the pancreas toward the lower left, exposing the splenic hilum. The surgeon separated the membrane of the body and tail of the pancreas to reach the posterior space at the superior border of the pancreas, and opened the vascular envelope at the end of the splenic arteries. The surgeon dissected away the lymphatic fatty tissue on the surface of the inferior splenic lobar artery from the lower pole of the spleen, vascularised the left gastroepiploic artery issuing from the inferior splenic lobar artery, and then cut the left gastroepiploic artery (No. 4sb) from the origin. Next, the assistant put the free omentum between the liver and the stomach and continually pulled the posterior wall of the fundus and body of the stomach to the upper right. The surgeon gently pressed the pancreas to fully reveal the retropancreatic space and the space inside the splenorenal ligament. Then, the surgeon tracked the termini of the splenic vessels along the completely vascularised the lower lobar vessels of the spleen within the space inside the splenorenal ligament. Next, the surgeon carefully dissected the fatty lymphatic tissue around the splenic vessels (No. 11d) along the latent anatomic spaces on the surface of the splenic vessels. At this time, the assistant gently pulled up the lymphatic fatty tissue at the surface of the inferior splenic lobar artery. Starting from the root of the left gastroepiploic artery, the surgeon, using the non-functional face of the ultrasonic scalpel, closed the surface of the inferior splenic lobar artery. The surgeon used the ultrasonic scalpel to carefully dissect the lymphatic fatty tissue and to vascularise the inferior splenic lobar artery. After the latter became visible, the short gastric arteries issuing from the inferior splenic lobar artery were skeletonised and divided at their roots, resulting in complete vascularisation of the inferior splenic lobar artery. The fatty tissues and gastric tissues were pulled up by the assistant, and the surgeon dissected the lymphatic fatty tissue on the surface of the superior splenic lobar artery, starting from the root of the artery towards the upper pole of the spleen, as described for vascularisation of the inferior splenic lobar artery. One branch of the short gastric artery issuing from the superior splenic lobar artery was skeletonised and divided at its root. This procedure resulted in LN dissections at the front of the splenic vessels (Figure 3). The assistant then pulled the root of the inferior splenic lobar artery towards the upper right, revealing the lymphatic fatty tissue behind the splenic hilum. The latter was pulled up by the surgeon towards the lower left to maintain tension. The lymphatic fatty tissue behind the splenic hilum was then dissected (No. 10) (Figure 4). A piece of gauze was placed behind the splenic hilum to indicate that the vessels had been vascularised and the LNs had been completely dissected.

**Digestive tract reconstruction**: The duodenum was transected 2 cm below the pylorus with a 60-mm laparoscopic cartridge linear stapling device through the major hand port. Finally, a longitudinal laparotomy was performed using a 6–8 cm skin incision at the epigastrium, and the specimen was extracted from the peritoneal cavity. The transaction of the esophagus and Roux-en-Y esophagojejunostomy was carried out using a circular stapler. A side-to-side jejunojejunostomy was performed by hand suture.

**RESULTS**

***Patient clinicopathological characteristics***

The 108 patients included 87 males (80.6%) and 21 females (19.4%) with a mean age of 62.5 ± 9.2 years (range, 24 to 82 years) and mean body mass index (BMI) of 22.1 ± 2.9 kg/m2 (range, 14.5 to 34.5 kg/m2). The preoperative clinical T stage was cT2 in 36 patients (33.3%) and cT3 in 72 patients (66.7%). The postoperative pathologic TNM stages included pT1 (*n* = 12), pT2 (*n* = 14), pT3 (*n* = 73), and pT4a (*n* = 9); pN0 (*n* = 31), pN1 (*n* = 17), pN2 (*n* = 23), and pN3 (*n* = 37); IA (*n* = 9), IB (*n* = 8), IIA (*n* = 19), IIB (*n* = 14), IIIA (*n* = 23), IIIB (*n* = 29), and IIIC (*n* = 6) (Table 1).

***Intraoperative and postoperative surgical outcomes***

For all 108 patients, the mean operation time was 169.3 ± 27.1 min, and the mean No. 10 lymphadenectomy time was 20.0 ± 5.7 min. The mean estimated blood loss was 46.2 ± 11.3 mL, and the mean estimated blood loss for No. 10 lymphadenectomy was 14.3 ± 3.8 mL. The mean times to first flatus, fluid diet, and soft diet were 3.4 ± 1.1, 4.7 ± 1.6, and 8.3 ± 4.2 d, respectively, and the mean postoperative hospital stay was 11.9 ± 6.0 d (Table 2).

***Morbidity and mortality***

Four patients experienced intraoperative complications; thus, the intraoperative morbidity rate was 3.7%. One patient experienced each of the following complications: injury to the transverse colon, injury to the splenic envelope, bleeding from the gastric coronary vein, and bleeding from the gastric short arteries. All complications were successfully treated during laparoscopic surgery. No patient required conversion to laparotomy, and no patient required splenectomy because of intraoperative injury to the splenic blood vessels or the spleen itself. Postoperative complications occurred in 13 patients; thus, the postoperative morbidity rate was 12.0%. These complications included abdominal infection in two patients, pulmonary infection in eight patients, inflammatory intestinal obstruction in one patient, chylous fistula in one patient, and anastomotic leakage in one patient. These postoperative complications were all successfully treated with conservative methods, and none of these patients required a second operation (Table 3). No patient experienced an operative splenic infarction, hemorrhage of the splenic blood vessels, or complications of spleen itself. The 30-d mortality rate for the total patient population was 0%.

***LN dissection***

The total number of LNs in all 108 patients was 4797, with a mean of 44.4 ± 17.6 LNs retrieved from each specimen. The total number of No. 10 LNs in all patients was 327, with a mean of 3.0 ± 2.4 No. 10 LNs retrieved per patient. Three patients (2.8%) had LN metastasis of the splenic hilus, including two patients with pT3 tumours and one patient with pT4a tumours, all located in the greater curvature (Table 4). There was no No. 10 LN metastasis in the patients with pT1 and pT2 tumours.

***Postoperative follow-up***

The 108 patients were followed up for a median 18 mo (range, 12 to 23 mo). No patient died or experienced tumour recurrence or metastasis during the follow-up period.

**DISCUSSION**

D2 lymphadenectomy, including the removal of No. 10 LNs, has become the standard surgical procedure for patients with curable AUGC[1,6]. In recent years, with the advances in surgical concepts, the improvements in the anatomical techniques and the progress of organ retention, spleen-preserving No. 10 LN dissection has been increasingly used for AUGC patients[2,7,9,12]. However, this procedure is technically difficult not only because of the intricate and complex blood vessels but also because of the deep and limited operative space in the splenic hilum. On one hand, in open surgery, the complete removal the No. 10 LNs often requires the mobilisation of the spleen from the abdominal cavity, which obviously increases patient trauma, elongates operation time, and is especially difficult for obese patients and patients with splenic adhesions. On the other hand, maintaining the spleen within the abdominal cavity and performing spleen-preserving No. 10 LN dissection directly would not completely remove all LNs because the exposure would be insufficient. Similarly with open surgery, spleen-preserving No. 10 LN dissection is also one of the most difficult procedures in laparoscopic surgery. Previously, a few studies have reported on the feasibility of laparoscopic spleen-preserving splenic hilar LN dissection for AUGC[17-19] patients , but the sample sizes in these studies was small, and the value of the procedure needed to be further evaluated by studies with large samples. According to the 3rd English edition of Japanese classification of gastric carcinoma[1], splenic hilar lymphadenectomy is unnecessary for cT1 tumors and laparoscopic surgery applied to cT4 tumors has been controversial. In the current study, therefore, we studied the feasibility and short-term efficacy of laparoscopic spleen-preserving No. 10 lymphadenectomy in consecutive 108 patients with stage cT2-T3 upper gastric cancer. Our data showed that the average time needed for No. 10 LN dissection was approximately 20 minutes, with less bleeding and shorter postoperative hospital stays, suggesting that laparoscopic spleen-preserving No. 10 lymphadenectomy is technically feasible.

Previous studies have reported that the intraoperative complication rate of laparoscopic gastric surgery was 2.6%-4.4%[23-24]. Consistent with these findings, we observed intraoperative complications in 4 of 108 patients(3.7%). None of our patients required conversion to laparotomy, and no patient required splenectomy because of injury to the spleen or splenic blood vessels. Postoperative complications have been reported in 8.7%-25.0% of patients who have undergone open spleen-preserving No. 10 lymphadenectomy for upper gastric cancer[2,9,11-12], and a recent study reported postoperative complications in 2 of 15 (13.3%) patients with upper gastric cancer who underwent laparoscopic spleen-preserving No. 10 lymphadenectomy[17]. In the current study, we found that 13 of 108 patients (12.0%) experienced postoperative complications, but no patient died within 30-d follow-up, suggesting that laparoscopic spleen-preserving No. 10 LNs dissection is safe and does not increase postoperative morbidity and mortality rates. In our experience, the keys to No. 10 LN dissection are a skilled laparoscopic technique, familiarity with the minimally invasive vascular anatomy of the splenic hilum area, and a cooperative surgical team. Moreover, the laparoscope, with its unique perspective, lighting and amplification, can more clearly visualise the splenic vasculature, nerves, fascia, and other structures, thereby reducing damage to the splenic vessels and spleen and assisting the surgeon in performing spleen-preserving No. 10 lymphadenectomy without splenic mobilisation.

The number of dissected LNs is an important assessment of the outcome of LN dissection. The average number of No. 10 LNs dissected per patient has been reported to be 3 LNs during open radical surgery for upper gastric cancer involving splenectomy[25] and 1.7 LNs during open radical surgery with spleen-preserving No. 10 lymphadenectomy[26]. During laparoscopic spleen-preserving No. 10 lymphadenectomy, the average numbers of No. 10 LNs dissected per patient were reported to be 2.7[17] and 2.6[19], indicating that a similar number of No. 10 LNs were dissected during laparoscopic and open surgery. In the current study, the average number of No. 10 LNs dissected was 3.0, which was similar to the other reports. No. 10 LNs are prone to metastasis in AUGC[25], and the metastasis rate to the No. 10 LNs reportedly ranges from 5.1% to 20.9%[9-11, 27-28]. Moreover, the No. 10 LNs metastasis rate is related to the tumour location, depth of invasion, other total LNs metastasis status and size of the primary tumour[9-11, 27-28]. In the current study, we observed metastases in these LNs in only 3 of 108 patients (2.8%), including two patients with pT3 and one patient with pT4a tumours, all located in the greater curvature, but there was no No. 10 LNs metastasis in the patients with pT1 and pT2 tumours. Therefore, this present study suggested that routine No. 10 lymphadenectomy may be unnecessary for AUGC without serosa invasion, unless T3 tumours are located in the greater curvature.

Patient survival after radical gastrectomy is important in evaluating its efficacy. The short and long term survival rates were greater in patients undergoing open spleen-preserving No. 10 lymphadenectomy for the treatment of upper stomach cancer[4-6]. In Hyung’s study, none of the 15 patients who underwent a laparoscopic procedure died or experienced tumour recurrence after a median follow-up period of 21 mo[17]. We found that after a median follow-up time of 18 mo, none of our patients experienced recurrence or metastasis. Longer follow-up periods, however, are required to determine the long-term efficacy of this procedure.

In conclusion, laparoscopic spleen-preserving No. 10 lymphadenectomy is feasible and effective for patients with AUGC. However, routine No. 10 lymphadenectomy may be unnecessary for AUGC without serosa invasion, unless T3 tumours are located in the greater curvature. In addition, multi-centre, prospective, randomised controlled studies involving greater numbers of patients and longer follow-up times, are needed to confirm its long-term efficacy.

**COMMENTS**

***Background***

Standard D2 lymph node (LN) dissection during total gastrectomy for advanced upper gastric cancer (AUGC) requires the removal of the No. 10 LNs, according to the 3rd English edition of Japanese classification of gastric carcinoma. Total gastrectomy with spleen-preserving No. 10 lymphadenectomy is increasingly used in open surgery to treat patients with upper gastric cancer. Several studies have shown that laparoscopic spleen-preserving No. 10 lymphadenectomy is feasible for patients with upper gastric cancer, but the sample sizes in these studies were small, and the value of the procedure must be further evaluated by studies with large sample sizes. Furthermore, it remains controversial whether routine No. 10 lymphadenectomy should be performed for AUGC without serosa invasion.

***Research frontiers***

Laparoscopic spleen-preserving No. 10 lymphadenectomy has been difficult to accomplish because of the possibilities of injury to splenic vessels and parenchyma of the spleen or pancreas.

***Innovations and breakthroughs***

The results of the current study demonstrate that the average time needed for laparoscopic spleen-preserving No. 10 lymphadenectomy was approximately 20 min and included less bleeding and shorter postoperative hospital stays. Moreover, the intraoperative and postoperative morbidity rates were 3.7% and 12.0%, respectively, but there was no postoperative mortality. At the same time, a mean of 3.0 ± 2.4 No. 10 LNs retrieved per patient. Three patients (2.8%) had LN metastasis of the splenic hilus, including two patients with pT3 and one patient with pT4a tumours, all located in the greater curvature. At the median follow-up of 18 mo (range, 12 to 23 mo), no patient died or experienced tumour recurrence or metastasis during the follow-up period.

***Applications***

The study results suggest that laparoscopic spleen-preserving No. 10 lymphadenectomy is feasible and effective for AUGC. Routine No. 10 lymphadenectomy may be unnecessary for AUGC without serosa invasion, unless T3 tumours are located in the greater curvature. The authors believe our results should encourage more surgeons to perform laparoscopic total gastrectomy with No. 10 lymphadenectomy and facilitate the acceptance of this procedure as a surgical option for AUGC patients.

***Terminology***

Spleen-preserving No. 10 lymphadenectomy, surgeons didn’t require to remove the spleen during the No. 10 lymphadenectomy when performing total gastrectomy with D2 LN dissection. Body mass index (BMI) was used as an objective index to indicate massive obesity. The cut-off value was chosen according to the World Health Organisation guidelines for the Western Pacific region.

***Peer review***

This is a good work in which the authors evaluate the feasibility and short-term efficacy of laparoscopic spleen-preserving No. 10 lymphadenectomy for AUGC. Congratulate the authors for the excellence of the work. All the contents in this study are appropriately presented. This manuscript is well written and documented. Additionally, this manuscript provides additional new knowledge to the literature.

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**P-Reviewer:** Abd Ellatif ME, Coskun A, Ferreira Caboclo JL

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**E:\2013-10-10\2014 整理稿件\Na Ma-2014-06-19-转苟苏鑫\8797\8797 figure\Figure 1.tif**

**Figure 1 Preoperative computed tomography angiography showing the drainage of the splenic arteries.** Abdominal aorta (arrow); splenic a (arrow). a: Artery.

E:\2013-10-10\2014 整理稿件\Na Ma-2014-06-19-转苟苏鑫\8797\8797 figure\Figure 2.tif

**Figure 2 Preoperative computed tomography angiography showing the drainage of the splenic veins.** Abdominal aorta (arrow); splenic a (arrow); splenic v (arrow).a: Artery; v: Vein.

E:\2013-10-10\2014 整理稿件\Na Ma-2014-06-19-转苟苏鑫\8797\8797 figure\Figure 3.tif

**Figure 3 No. 10 lymph nodes lymphadenectomy at the front of the splenic vessels (**[**anterior**](http://dict.youdao.com/w/anterior/) **view).** Dividing left gastroepiploic a (arrow); Dividing short gastric a (arrow);splenic a (arrow); splenic v (arrow); a: Artery; v: Vein.

E:\2013-10-10\2014 整理稿件\Na Ma-2014-06-19-转苟苏鑫\8797\8797 figure\Figure 4.tif

**Figure 4 No. 10 lymph nodes lymphadenectomy behind the splenic vessels (posterior view).** Dividing left gastroepiploic a (arrow); splenic vein (arrow); a: Artery; v: Vein.

**Table 1 Baseline patient demographic and clinicopathological characteristics**

|  |  |  |
| --- | --- | --- |
| **Characteristics** |  | **Value** |
| Gender | Male/female | 87/21 |
| Age (yr) |  | 62 ± 9 |
| Tumour size (cm) |  | 5.0 ± 2.6 |
| BMI (kg/m2) |  | 22.1(14.5-34.5) |
| Tumour location | Lesser curvature/greater curvature/anterior wall /posterior wall/circumferential involvement | 30/21/15/19/23 |
| Pathological type | Differentiated/undifferentiated type | 43/65 |
| cT stage | T2/T3 | 36/72 |
| pT stage | T1/T2/T3/T4a | 12/14/73/9 |
| pN stage | N0/N1/N2/N3 | 31/17/23/37 |
| TNM stage | IA/IB/IIA/IIB/IIIA/IIIB/IIIC | 9/8/19/14/23/29/6 |

Data are expressed as mean ± SD. BMI: Body mass index; cT stage: Clinical tumor stage; pT stage: Pathological tumor stage.

**Table 2 Intraoperative and postoperative surgical outcomes**

|  |  |
| --- | --- |
| **Items** | **Value** |
| Operation time (min) | 169.3 ± 27.1 |
| Blood loss (mL) | 46.2 ± 11.3 |
| No. 10 lymphadenectomy (min) | 20.0 ± 5.7 |
| No. 10 lymphadenectomy blood loss (mL) | 14.3 ± 3.8 |
| Time to first flatus (POD) | 3.4 ± 1.1 |
| Time to fluid diet (POD) | 4.7 ± 1.6 |
| Time to soft diet (POD) | 8.3 ± 4.2 |
| Hospital stay (POD) | 11.9 ± 6.0 |

Data are expressed as mean ± SD. No. 10 lymphadenectomy: Splenic hilar lymphadenectomy; POD: Postoperative days.

**Table 3 Intraoperative and postoperative complications**

|  |  |  |
| --- | --- | --- |
| **Items** | **Value** | **Incidence rate** |
| Intraoperative complications (*n*) | 4 | 3.7% |
| Transverse colon injury | 1 |  |
| Spleen injury | 1 |  |
| Left gastric vein bleeding | 1 |  |
| Gastric short arteries bleeding | 1 |  |
| Postoperative complications (*n*) | 13 | 12.0% |
| Pulmonary infection | 8 |  |
| Abdominal infection | 2 |  |
| Anastomotic leakage | 1 |  |
| Intestinal obstruction | 1 |  |
| chylous fistula | 1 |  |

**Table 4 Lymph nodes dissection results**

|  |  |
| --- | --- |
| **Items** | **Value** |
| Total No. of retrieved LNs | 4797 |
| Mean No. of retrieved LNs | 44.4 ± 17.6 |
| Total No. of retrieved No. 10 LNs | 327 |
| Mean No. of retrieved No. 10 LNs | 3.0 ± 2.4 |
| Total No. of No. 10 LNs metastasis | 3 |
| No. of No. 10 LNs [metastasis](http://dict.youdao.com/w/metastasis/) in pT3 | 2 |
| No. of No. 10 LNs [metastasis](http://dict.youdao.com/w/metastasis/) in pT4a | 1 |
| No. 10 LNs [metastasis](http://dict.youdao.com/w/metastasis/) rate(%) | 2.8 |
| No. 10 LNs [metastasis](http://dict.youdao.com/w/metastasis/) rate in pT3(%) | 2.7 |
| No. 10 LNs [metastasis](http://dict.youdao.com/w/metastasis/) rate in pT4a(%) | 11.1 |

Data are expressed as mean ± SD. LN: Lymph node; No. 10 LN: Splenic hilar lymph node; pT3: Pathological T3 stage; pT4a: Pathological T4a stage.