Ligation of left gastric vein may cause delayed gastric emptying after pancreateoduodenectomy: a retrospective study

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Abstract

Background: This study aimed to determine which running pattern of the left gastric vein (LGV) is most frequently ligated in subtotal stomach-preserving pancreateoduodenectomy (SSPPD) and how LGV ligation affects delayed gastric emptying (DGE) after SSPPD.

Methods: We retrospectively analysed 105 patients who underwent SSPPD between January 2016 and September 2021. We classified the running pattern of LGV as follows: type 1 runs dorsal to the common hepatic artery (CHA) or splenic artery (SpA) to join the portal vein (PV), type 2 runs dorsal to the CHA or SpA and joins the splenic vein, type 3 runs ventral to the CHA or SpA and joins the PV, and type 4 runs ventral to the CHA or SpA and joins the SpV. Univariate and multivariate analyses were used to identify differences between patients with and without DGE after SSPPD.

Results: Type 1 LGV running pattern was observed in 47 cases (44.8%), type 2 in 23 (21.9%), type 3 in 12 (11.4%), and type 4 in 23 (21.9%). The ligation rate was significantly higher in type 3 (75.0%) LGVs (p < 0.0001). Preoperative obstructive jaundice (p = 0.0306), LGV ligation (p < 0.0001), grade B or C pancreatic fistula (p = 0.0116), and sepsis (p = 0.0123) were risk factors for DGE in the univariate analysis. Multivariate analysis showed that LGV ligation was an independent risk factor for DGE (odds ratio: 13.60, 95% confidence interval: 3.80–48.68, p < 0.0001).

Conclusion: Type 3 LGVs are often ligated because they impede lymph node dissection; however, LGV preservation may reduce the occurrence of DGE after SSPPD.

Keywords: Obstructive jaundice, Pancreatic fistula, Risk factor, Sepsis, Stomach

Background

Pancreateoduodenectomy (PD) is the standard surgery for tumours in the pancreatic head, distal common bile duct, and duodenal papilla [1]. Despite the recent establishment and improvement of various operative techniques and perioperative management protocols, PD still results in several postoperative complications [2, 3].

Delayed gastric emptying (DGE) is a complication of PD, occurring in 19–57% of cases [4]. Although DGE is not life-threatening, it exacerbates patients’ quality of life and prolongs hospitalisation after surgery [5]. Surgeons have cogitated how to prevent the development of DGE after PD. Several studies have reported that pylorus-preserving PD (PPPD) [6], ischaemic anastomotic sites of alimentary tract reconstruction,
and injury of the branches of the vagus nerve [4] could be risk factors for the development of DGE after PD. Moreover, intra-abdominal infection and postoperative pancreatic fistula (POPF) are well-known risk factors for DGE after PD [7]. In addition, the retrocolic route for alimentary tract reconstruction [1], absence of Braun enterenterostomy [8], and Roux-en-Y reconstruction method [9] have been reported as risk factors for DGE after PD.

However, little attention has been focused on gastric venous congestion following ligation of the coronary vein, which consists of the LGV and right gastric vein. Surgeons often encounter situations where the LGV requires ligation during lymph node dissection around the PV and/or CHA. The LGV occasionally obstructs the surgical field of vision in lymph node dissection and can be injured and cause bleeding or impede the surgery through other factors. Kuroasaki et al. reported that preservation of the LGV reduces the occurrence of DGE after PPPD [10]. They suggested that LGV ligation induces antroduodenal congestion after PPPD. However, the effect of LGV ligation on DGE after subtotal stomach-preserving pancreaticoduodenectomy (SSPPD), in which the pyloric ring is excised, remains unclear.

In this study, we classified the original running pattern of the LGV and investigated which type of LGV was most frequently ligated. Furthermore, we examined the effect of LGV ligation on DGE after SSPPDs.

Methods
In this study, we retrospectively investigated the running pattern of the LGV and development of DGE after ligation of the LGV in SSPPD.

Patients
A total of 105 adults (>20 years old) who underwent SSPPD for pancreatic head tumours, periampullary tumours, or for diseases manifesting as a pancreatic mass at Matsuyama Red Cross Hospital between January 2016 and September 2021 were included in this study. Patients scheduled to undergo total pancreatectomy, combined liver resection, and pancreaticogastrostomy and those with a history of gastric surgery or colorectal surgery were excluded. All treatment procedures were performed after informed consent was obtained from the patients. Medical charts were retrospectively reviewed to obtain patient data. This study was approved by the Clinical Study Examination Committee of the Matsuyama Red Cross Hospital (Approval No. 919) and was performed in accordance with the ethical standards of the 1964 Declaration of Helsinki and its subsequent amendments.

Surgical procedures
SSPPD with lymph node resection was performed as a standard procedure for pancreatic head, distal common bile duct, or periampullary tumours. The SSPPD involved a division of the stomach 3 cm proximal to the pyloric ring, followed by resection of the entire duodenum, gallbladder, distal common bile duct, and pancreatic head.

We resected the right gastric artery and vein for routine dissection of #5 lymph node for all surgical procedures in SSPPD [11]. Simultaneously, we performed #12 and #8 lymph node dissections and resection of the gastroduodenal artery. The inferior pancreaticoduodenal artery was resected routinely. The lymph nodes of the right semicircle of the superior mesenteric artery were dissected (#14 lymph node dissection) for pancreatic carcinomas.

Reconstruction was performed using the modified Child method. Either Roux-en-Y or Billroth II reconstruction was performed using the modified Child method. Pancreaticojejunostomy was performed using the modified Blumgart anastomosis [12] or Kakita method [13]. There were no restrictions on the hepaticojejunostomy technique used. Alimentary tract reconstruction as side-to-side gastroenterostomy was performed by using an automatic stapling device or hand-sewn anastomosis. Braun anastomosis was performed in all patients with Billroth II reconstruction.

Antecolic alimentary tract reconstruction was performed in all cases. During antecolic reconstruction, the anastomosis was positioned anterior to the transverse colon. The number, types, and locations of the intra-abdominal drainage tubes were determined according to the surgeon’s preference.

Postoperative management
The patients were administered epidural anaesthesia for 3 days, as appropriate. Early mobilisation was encouraged. The nasogastric tube (NGT) was routinely removed on the morning of postoperative day (POD) 1 if the drainage volume was <200 mL. If a patient vomited persistently or the drainage volume was >200 mL, an NGT was maintained. Conventionally, a solid diet was initiated on POD 3. However, this schedule was changed according to clinical observations, such as abdominal swelling, little peristaltic sounds, or vomiting. Drainage fluid biochemistry was measured for the presence of postoperative bile leakage or POPF by countering bilirubin and amylase on PODs 1, 3, 5, and 7. Drains were removed if there were no signs of postoperative biliary leakage, POPF, or chylorrhea. Postoperatively, a proton pump inhibitor was administered orally from POD 1 to prevent gastrointestinal ulceration. Octreotide administration was determined according to the attending surgeon’s preferences.
Pancreatic enzyme supplementation, mosapride citrate hydrate, or Japanese herbal medicine daikenchuto was administered according to the patient’s condition.

**Postoperative complication definition**

**DGE**

DGE was defined and graded according to the International Study Group of Pancreatic Surgery consensus definition published in 2007 as follows: Grade A, NGT insertion after POD 3 or the inability to tolerate intake of solid diet by POD 7; Grade B, NGT required for 8–14 days postoperatively, NGT reinsertion after POD 7, or the inability to tolerate solid diet by POD 14; and Grade C, NGT required for >14 days postoperatively, NGT reinsertion after POD 14, or the inability to tolerate solid diet by POD 21 [4].

**POPF**

According to the 2016 update of the International Study Group for Pancreatic Fistula definition and grading of POPF, POPF is defined as a drain output amylase level >3 times the upper limit of the institutional normal serum amylase activity. Grade A POPF is defined and termed as a "biochemical leak", because it has no clinical importance and is no longer referred to as a true pancreatic fistula. Grade B requires a change in postoperative management; drains are either left in place for >3 weeks or repositioned through endoscopic or percutaneous procedures. Grade C refers to POPFs that require reoperation or lead to single or multiple organ failure and/or mortality attributable to pancreatic fistula [14].

**Other complications**

Postoperative intra-abdominal infection was defined as the verification of a positive bacterial culture from intra-abdominal drainage tubes after surgery. Sepsis was defined as the detection of bacteria and/or fungi in blood cultures.

**Running pattern of LGV classification**

Running pattern of LGV was detected by using computed tomography performed before and after surgery and intraoperative findings from operation records. We performed computed tomography on seven days from surgery routinely. Then, we have investigated that the LGV was preserved or not in postoperative computed tomography. Moreover, we have added presence of LGV ligation in operation record. We investigated the anatomy of the LGV and classified it into four types according to the running pattern of the LGV, ventral or distal route of the common hepatic artery (CHA) and splenic artery (SpA) arcade, and joining point to the veins of the portal vein (PV) or splenic vein (SpV). Previous reports have identified three positions for LGV confluence into major veins: the PV, SpV, and junction of the PV and SpV [15, 16]. However, Kawasaki et al. described a relationship between the LGV and arteries around the pancreas [17]. In this study, we classified the running pattern of the LGV to simplify the classification from a surgical and procedural viewpoint.

Type 1 LGV runs dorsal to the CHA or SpA (mostly dorsal to the CHA) and join the PV. Type 2 LGV runs dorsal to the CHA or SpA (mostly dorsal to the SpA) and joins the SpV. Type 3 LGV runs ventral to the CHA or SpA (mostly ventral of CHA) and joins the PV. Finally, type 4 LGV runs ventral to the CHA or SpA (mostly ventral to the SpA) and joins the SpV (Fig. 1). LGV ligation data were collected from operation records.

We found two cases which LGV flows into the left branch of the portal vein. They were excluded in this study.

**Statistical analyses**

All values are expressed as mean and standard deviation. Categorical variables were compared using χ² tests. Statistical significance was set at \( p < 0.05 \). A logistic regression analysis was used for the multivariate analysis of

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**Fig. 1** Four running patterns of the left gastric vein. Yellow and green arrowheads point to the left gastric veins. The running patterns indicated by the yellow arrowheads were more frequently ligated than the patterns indicated with green arrowheads. LGV, left gastric vein; CHA, common hepatic artery; SpA, splenic artery; DGA, duodenal gastric artery; PHA, proper hepatic artery; PV, portal vein; SpV, splenic vein; SMV, superior mesenteric vein.
Results

Patient characteristics

A total of 105 patients underwent SSPPD for pancreatic head tumours, periampullary tumours, or diseases manifesting as pancreatic masses between January 2016 and September 2021. The patient characteristics are described in Additional File 1. The most common diagnosis was pancreatic adenocarcinoma (54.3%).

Frequency of running pattern of LGV and LGV ligation

We investigated the running pattern of the LGV in all the patients and found that 47 cases (44.8%) were type 1, 23 (21.9%) were type 2, 12 (11.4%) were type 3, and 23 (21.9%) were type 4. We also examined which type of running pattern of the LGV was most frequently ligated. The number of ligated LGVs according to type was as follows: type 1, six cases (12.8%); type 2, one case (4.4%); type 3, nine cases (75.0%); and type 4, three cases (13.0%) (Table 1).

Intraoperative findings

Intraoperative findings are shown in Table 2. In this study, all surgical procedures were performed using the SSPPD. Reconstruction techniques (modified Child method), reconstruction method (Roux-en-Y or Billroth II), and alimentary tract reconstruction method (automatic stapling device or hand-sewn anastomosis) were determined according to the operating surgeon's preferences. In 19 patients (18.1%), the LGV was ligated during surgery.

Surgical outcomes

Postoperative factors are shown in Table 3. The frequency of all DGE grades after surgery was 20.0%. The frequency of grade B or C pancreatic fistulas was 33.3%. Wound infection occurred in 9.5% of patients and intra-abdominal infection developed in 22.9% of patients after surgery. Only one case of mortality occurred and was caused by disseminated intravascular coagulation after surgery. The deceased patient's right hepatic artery had to be resected and reconstructed because of cancer invasion. The reconstructed right hepatic artery did not develop occlusion but indicated stenosis, which led to the development of multiple liver abscesses and disseminated intravascular coagulation. The patient's general condition worsened, and he was unable to tolerate surgical or non-surgical treatment.

Univariate and multivariate analysis between DGE and non-DGE cases

Thereafter, we investigated the risk factors for DGE after surgery. The prevalence of preoperative obstructive jaundice was significantly higher in the DGE group than in the non-DGE group (66.7% vs. 40.5%, \( p = 0.0306 \)). Only LGV ligation showed a significant difference between the DGE and non-DGE groups (57.1% vs. 8.3%, \( p < 0.0001 \)) in intraoperative factors. On comparing the DGE and non-DGE groups, grade B or C pancreatic fistula (57.1% vs. 27.4%, \( p = 0.0116 \)) and sepsis (28.6% vs. 7.1%, \( p = 0.0123 \)) emerged as risk factors for the development of DGE after surgery. Postoperative hospitalisation was significantly longer in the DGE group than in the non-DGE group.

We subjected the risk factors for DGE revealed in the univariate analysis to a multivariate analysis and found that LGV ligation was the only independent risk factor for DGE after SSPPD (odds ratio: 13.60, 95% CI: 3.80–48.68, \( p < 0.0001 \)) (Table 4).

Univariate and multivariate analysis between DGE and non-DGE cases in LGV ligation group

Moreover, we also investigated the risk factors for DGE after surgery in LGV ligation group. On comparing the DGE and non-DGE groups, intra-abdominal infection (83.3% vs. 16.7%, \( p = 0.0051 \)) and sepsis (41.7% vs. 0.0%, \( p = 0.0258 \)) emerged as risk factors for the development of DGE after surgery in LGV ligation group. Postoperative hospitalisation was significantly longer in the DGE group than in the non-DGE group in LGV ligation group.

We subjected the risk factors for DGE in LGV ligation group revealed in the univariate analysis to a multivariate analysis and found that intra-abdominal infection was the only independent risk factor for DGE after SSPPD in LGV ligation group (odds ratio: 12.5, 95% CI: 0.84–186.3, \( p = 0.0413 \)) (Table 5).

DGE development with or without LGV ligation in the absence of grade B or C pancreatic fistula group

Furthermore, we investigated DGE development after surgery with or without LGV ligation in the absence of grade B or C pancreatic fistula. DGE was significantly

| Table 1 | Running pattern of LGV frequency and association between running patterns of the left gastric vein and ligation rate |
|---------|---------------------------------------------------------------|
| Running pattern of LGV | Patients (%) | Ligation (%) |
| Type 1 | 47 (44.8) | 6 (12.8) |
| Type 2 | 23 (21.9) | 1 (4.4) |
| Type 3 | 12 (11.4) | 9 (75.0) |
| Type 4 | 23 (21.9) | 3 (13.0) |
more frequent in the LGV ligation group than in the LGV preservation group (33.3% vs. 10.3%, \( p = 0.0493 \)).

**Discussion**

DGE is a major complication of PD; however, the development and risk factors of DGE are still controversial [1, 4, 5, 7–9]. Physiologically, the causes of DGE after PD are hypothesised as follows: decreased plasma levels of the hormone motilin due to resection of the duodenum and its M-cells [18, 19], damage to branches of the vagus nerve in a dissection of lymph nodes, and ischaemia of the point of the alimentary tract reconstruction [4]. DGE in the postoperative period decreases the quality of life of patients and prolongs hospitalisation by extension of inanition states, ultimately leading to increased health care costs [5]. In this study, DGE after SSPPD was more frequent in the LGV ligation group than in the LGV preservation group. We resected the right gastric artery and vein for routine dissection of #5 lymph nodes in performing SSPPD. LGV ligation might cause stasis of the lesser curvature of the stomach and lead to the development of DGE after SSPPD. Thus, LGV preservation may reduce the rate of DGE development after SSPPD (Fig. 2).

Moreover, LGVs with type 3 running pattern were the most frequently ligated LGV type. A type 3 LGV running ventral to the CHA or SpA and directly joining the PV would obstruct the surgical field of view in the dissection of #12p and/or #8a lymph nodes around the PV and CHA and could be ligated during lymph node dissection. Contrastingly, LGVs with type 1, type 2, and type 4 running patterns remain distant from the lymph node dissection field in SSPPD and thus are less likely to be ligated intraoperatively. It would be essential to preserve the LGV by taping it if it is recognised during surgery.
Several reports have revealed that intra- and postoperative management techniques reduce the incidence of DGE after surgery. Matsumoto et al. reported that the occurrence rate of DGE after surgery was higher in the PPPD group than in the SSPPD group [20]. Postoperative intra-abdominal infection and POPF are well-known risk factors for DGE after PD [7]. Tani et al. first performed a randomised controlled trial to investigate the effect of alimentary tract reconstruction (antecolic or retrocolic route) on DGE after PD [21]. Their results revealed a significantly higher occurrence rate of DGE after PPPD with the retrocolic alimentary route than with the antecolic route. In addition, Toyama et al. failed to prove the non-inferiority of retrocolic alimentary tract reconstruction to antecolic alimentary tract reconstruction with respect to the postoperative incidence of DGE [1]. Moreover, Hwang et al. suggested that Braun anastomosis reduced DGE after PD [22], and Shimoda et al. reported that the incidence of DGE after SSPPD can be decreased using

### Table 3 Postoperative factors

| Factors                        | Patients (n = 105) |
|-------------------------------|-------------------|
| Delayed gastric emptying (%)  |                   |
| None                          | 84 (80.0)         |
| Grade A                       | 14 (13.3)         |
| Grade B                       | 5 (4.8)           |
| Grade C                       | 2 (1.9)           |
| Pancreatic fistula (%)        |                   |
| None                          | 54 (51.4)         |
| Grade A                       | 16 (15.2)         |
| Grade B                       | 33 (31.4)         |
| Grade C                       | 2 (1.9)           |
| Wound infection (yes, %)      | 10 (9.5)          |
| Intra-abdominal infection (yes, %) | 24 (22.9) |
| Sepsis (yes, %)               | 12 (11.4)         |
| > 30 days mortality (yes, %)  | 1 (1.0)           |
| Postoperative hospitalisation (days), mean±standard deviation | 24.2 ± 14.5 |

### Table 4 Univariate and multivariate analysis of the association between delayed gastric emptying and operative factors

| Factors                        | Univariate | Multivariate |
|-------------------------------|------------|--------------|
|                               | Yes (n = 21) | No (n = 84) | P-value | Odds ratio (95% CI) | P-value |
| Sex (male, %)                 | 13 (61.9)  | 51 (60.7)   | 0.9203  |                     |         |
| Age (years), mean±standard deviation | 71 ± 8   | 70 ± 10    | 0.7734  |                     |         |
| BMI (kg/m²), mean±standard deviation | 22.1 ± 2.9 | 20.9 ± 3.6 | 0.1746  |                     |         |
| Diabetes mellitus (yes, %)    | 5 (23.8)   | 33 (39.3)  | 0.1759  |                     |         |
| Current smoker (yes, %)       | 2 (9.5)    | 14 (16.7)  | 0.3933  |                     |         |
| Preoperative obstructive jaundice (Yes, %) | 14 (66.7) | 34 (40.5) | 0.0306  | 3.13 (0.91–10.82)   | 0.0710  |
| Tumor size (mm), mean±standard deviation | 24.8 ± 2.6 | 26.2 ± 1.3 | 0.6037  |                     |         |
| Blood vessels invasion (yes, %) | 3 (14.9)  | 19 (22.6)  | 0.3848  |                     |         |
| Malignancy (yes, %)           | 18 (85.7)  | 69 (82.1)  | 0.6925  |                     |         |
| Pancreatic gland texture (hard, %) | 7 (33.3)  | 37 (44.1)  | 0.3689  |                     |         |
| Portal vein resection (yes, %) | 3 (14.3)  | 16 (19.1)  | 0.7895  |                     |         |
| Blood loss (g), mean±standard deviation | 368 ± 243 | 507 ± 407  | 0.1354  |                     |         |
| Intraoperative transfusion (yes, %) | 2 (9.5)   | 10 (11.9)  | 0.6383  |                     |         |
| Operative time (min), mean±standard deviation | 498 ± 109 | 534 ± 104 | 0.1574  |                     |         |
| Running pattern of LGV         | 11 (52.4)  | 36 (42.9)  | 0.0011  |                     |         |
| Type 1                        |            |             |         |                     |         |
| Type 2                        | 0 (0.0)    | 23 (27.4)  |         |                     |         |
| Type 3                        | 6 (28.6)   | 6 (7.1)    |         |                     |         |
| Type 4                        | 4 (19.1)   | 19 (22.6)  |         |                     |         |
| LGV ligation (yes, %)         | 12 (57.1)  | 7 (8.3)    | <0.0001 | 13.60 (3.80–48.68)  | <0.0001 |
| Stapling device for gastroenterostomy (yes, %) | 18 (85.7) | 77 (91.7) | 0.4273  |                     |         |
| Roux-en-Y reconstruction (yes, %) | 8 (38.1) | 47 (56.0) | 0.1428  |                     |         |
| Grade B or C pancreatic fistula (yes, %) | 12 (57.1) | 23 (27.4) | 0.0116  | 1.70 (0.32–9.17)   | 0.5357  |
| Wound infection (yes, %)      | 4 (19.1)   | 6 (7.1)    | 0.1242  |                     |         |
| Intra-abdominal infection (yes, %) | 8 (38.1) | 16 (19.1) | 0.0749  |                     |         |
| Sepsis (yes, %)               | 6 (28.6)   | 6 (7.1)    | 0.0123  | 1.75 (0.31–9.80)   | 0.5231  |
| Postoperative hospitalisation (days), mean±standard deviation | 33 ± 12   | 22 ± 14    | 0.0017  |                     |         |
Billroth II rather than Roux-en-Y reconstruction for gastrojejunalostomy [9].

First, all cases in this study were SSPPD; therefore, the effect of PPPD on the development of DGE after surgery could not be determined. We propose that the mechanism of DGE development between SSPPDs and PPPDs should be dissociated with or without resection of the pyloric ring. In our study, there was no significant difference in the data between patients with and without intra-abdominal infection. We also examined the effect of LGV ligation for DGE after SSPPD in patients without grade B or C POPF. The LGV ligation group showed a significantly higher incidence of DGE after SSPPD than the LGV preservation group. This result suggests that LGV ligation as a risk factor for DGE after SSPPD is unrelated to the presence of POPF. The effect of the alimentary tract reconstruction route (antecolic or retrocolic route) on DGE was also unclear in this study because antecolic route reconstruction was performed in all cases. Braun anastomosis was routinely performed in all Billroth II reconstruction cases. Moreover, we investigated the effect of LGV ligation on DGE after SSPPD in patients who underwent Roux-en-Y reconstruction. The LGV ligation group showed a significantly higher rate of DGE incidence after SSPPD than the LGV preservation group (45.5% vs. 6.8%, \( p = 0.0012 \)). We also examined the effect of LGV ligation on DGE after surgery in patients who underwent Billroth II reconstruction. The LGV ligation group showed a significantly higher rate of DGE incidence after SSPPD than the LGV preservation group (87.5% vs. 14.3%, \( p = 0.0012 \)). These findings revealed that LGV ligation was a risk factor for DGE after SSPPD but was unrelated to the reconstruction methods.

### Table 5 Univariate and multivariate analysis of the association between delayed gastric emptying and operative factors in LGV ligation group

| Factors                                         | Univariate                | Multivariate                |
|------------------------------------------------|---------------------------|-----------------------------|
|                                                | DGE                       |                            |
|                                                | Yes (n = 12) | No (n = 6) | \( P \)-value | Odds ratio (95% CI) | \( P \)-value |
| Sex (male, %)                                   | 10 (83.3) | 3 (50.0) | 0.1436 |                       |              |
| Age (years), mean ± standard deviation          | 71 ± 3 | 69 ± 4 | 0.6748 |                       |              |
| BMI (kg/m²), mean ± standard deviation          | 22.0 ± 1.0 | 21.9 ± 1.4 | 0.9534 |                       |              |
| Diabetes mellitus (yes, %)                      | 1 (8.3) | 2 (33.3) | 0.1926 |                       |              |
| Current smoker (yes, %)                         | 1 (8.3) | 0 (0.0) | 0.3594 |                       |              |
| Preoperative obstructive jaundice (Yes, %)      | 8 (66.7) | 2 (33.3) | 0.1778 |                       |              |
| Tumor size (mm), mean ± standard deviation      | 27.2 ± 3.0 | 22.3 ± 3.9 | 0.3391 |                       |              |
| Blood vessels invasion (yes, %)                 | 0 (0.0) | 0 (0.0) |              |                       |              |
| Malignancy (yes, %)                             | 11 (91.7) | 4 (66.7) | 0.0777 |                       |              |
| Pancreatic gland texture (hard, %)              | 2 (16.7) | 2 (33.3) | 0.4319 |                       |              |
| Portal vein resection (yes, %)                  | 0 (0.0) | 0 (0.0) |              |                       |              |
| Blood loss (g), mean ± standard deviation        | 335 ± 250 | 524 ± 463 | 0.2705 |                       |              |
| Intraoperative transfusion (yes, %)              | 1 (8.3) | 1 (16.7) | 0.6052 |                       |              |
| Operative time (min), mean ± standard deviation | 476 ± 128 | 523 ± 60 | 0.4096 |                       |              |
| Running pattern of LGV                          | 0.2284 |              |              |                       |              |
| Type 1                                          | 5 (41.7) | 1 (16.7) |              |                       |              |
| Type 2                                          | 0 (0.0) | 1 (16.7) |              |                       |              |
| Type 3                                          | 6 (50.0) | 3 (50.0) |              |                       |              |
| Type 4                                          | 1 (8.3) | 2 (33.3) |              |                       |              |
| Stapling device for gastroenterostomy (yes, %)   | 9 (75.0) | 6 (100.0) | 0.0988 |                       |              |
| Roux-en-Y reconstruction (yes, %)               | 7 (58.3) | 1 (16.7) | 0.0821 |                       |              |
| Grade B or C pancreatic fistula (yes, %)        | 9 (75.0) | 1 (16.7) | 0.0861 |                       |              |
| Wound infection (yes, %)                        | 4 (33.3) | 0 (0.0) | 0.0515 |                       |              |
| Intra-abdominal infection (yes, %)              | 10 (83.3) | 1 (16.7) | 0.0051 | 12.5 (0.84–186.3) | 0.0413 |
| Sepsis (yes, %)                                 | 5 (41.7) | 0 (0.0) | 0.0258 | 1.69 (0.35–12.8) | 0.2551 |
| Postoperative hospitalisation (days), mean ± standard deviation | 36 ± 11 | 16 ± 7 | 0.0009 |                       |              |
Regarding the possibility of damage to the vagus nerve in DGE after SSPPD, Xu et al. suggested that preservation of the hepatic branches of the vagus nerve reduces the development of DGE after laparoscopic PD [23]. Nevertheless, the LGV runs closer to the caudal side of the hepatic branch of the vagus nerve, indicating that LGV ligation might be related to the damage of the left gastric artery plexus from the posterior trunk of the vagus nerve anatomically [24]. LGVs running ventral to the artery are easily ligated during lymph node dissection. Therefore, we suggest the possibility of injury to the left gastric artery plexus during LGV ligation. DGE is also a complication of distal gastrectomy [25]. Kim et al. demonstrated that the preservation of the proximal side of the bifurcation of the hepatic branch from the anterior trunk of the vagus nerve and celiac branch from the posterior trunk of the vagus nerve improves gastric function after distal gastrectomy [26]. We were unable to corroborate this observation in our study because we did not explore the damage to the vagus nerve branches; nevertheless, more popularisation of the laparoscopic PD procedure would enable us to visualise and investigate the running patterns of the thin vagus nerve branches in more detail.

This study has some limitations. First, we selected patients from a single centre. A multicentre study with a larger number of patients and greater variation in surgical techniques and surgeons would help us reach more definitive conclusions. Second, this was a retrospective study, and may have been subject to investigative bias. Further randomised controlled multicentre studies are needed.

**Conclusions**

We investigated variations in the running pattern of LGV and effect of LGV ligation on the occurrence rate of DGE after SSPPD. We found that type 3 LGVs running ventral to the CHA or SpA and joining the PV were most frequently ligated, and LGV ligation was an independent risk factor for DGE after SSPPD. Type 3 LGVs impede lymph node dissection around the PV and/or CHA and must be ligated. If the LGV is recognised during surgery, it should be preserved, as far as possible, by taping with vessel tapes. Since preservation of the LGV maintains gastric drainage venous flow, we suggest that the LGV should be preserved as far as possible to reduce gastric stasis and prevent the development of DGE after SSPPD.

**Abbreviations**

LGV: Left gastric vein; SSPPD: Subtotal stomach-preserving pancreatoduodenectomy; DGE: Delayed gastric emptying; CHA: Common hepatic artery; SpA: Splenic artery; PV: Portal vein; PD: Pancreatoduodenectomy; POPF: Postoperative pancreatic fistula; PPPD: Pylorus-preserving pancreatoduodenectomy; POD: Postoperative day; NGT: Nasogastric tube; SpV: Splenic vein.

**Supplementary Information**

The online version contains supplementary material available at https://doi.org/10.1186/s12876-022-02478-5.

**Additional file 1** Patient characteristics

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None

**Author contributions**

KK drafted the manuscript and formulated the study design and concepts. RM, KM, and TN critically revised the manuscript. TI, AO, TA, GK, DT, HY, and YK performed data collection. All authors have read and approved the final manuscript.

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**Availability of data and materials**

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

**Declarations**

**Ethics approval and consent to participate**

All treatment procedures were performed after informed consent was obtained from the patients. Medical charts were retrospectively reviewed to obtain patient data. This study was approved by the Clinical Study Examination Committee of the Matsuyama Red Cross Hospital (Approval No. 919) and was performed in accordance with the ethical standards of the 1964 Declaration of Helsinki and its subsequent amendments.

**Consent for publication**

Not applicable.
Competing interests
The author declare that they have no competing interests.

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