Systematic review of the incidence, presentation and management of gastroduodenal artery pseudoaneurysm after pancreatic resection

B. Brodie and H. M. Kocher

Background: Gastroduodenal artery (GDA) pseudoaneurysm is a serious complication following pancreatic resection, associated with high morbidity and mortality rates. This review aimed to report the incidence of GDA pseudoaneurysm after pancreatic surgery, and describe clinical presentation and management.

Methods: MEDLINE and Embase were searched systematically for clinical studies evaluating postoperative GDA pseudoaneurysm. Incidence was calculated by dividing total number of GDA pseudoaneurysms by the total number of pancreatic operations. Additional qualitative data related to GDA pseudoaneurysm presentation and management following pancreatic resection were extracted and reviewed from individual reports.

Results: Nine studies were selected for systematic review involving 4227 pancreatic operations with 55 GDA pseudoaneurysms, with a reported incidence of 1.3 (range 0.2–8.3) per cent. Additional data were extracted from 39 individual examples of GDA pseudoaneurysm from 14 studies. The median time for haemorrhage after surgery was at 15 (range 4–210) days. A preceding complication in the postoperative period was documented in four of 21 patients (67 per cent), and sentinel bleeding was observed in 14 of 20 patients (70 per cent). Postoperative complications after pseudoaneurysm management occurred in two-thirds of the patients (14 of 21). The overall survival rate was 85 per cent (33 of 39).

Conclusion: GDA pseudoaneurysm is a rare yet serious cause of haemorrhage after pancreatic surgery, with high mortality. The majority of the patients had a preceding complication. Sentinel bleeding was an important clinical indicator.

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Introduction

Mortality from pancreatic resection has fallen significantly over the past few decades, especially in experienced centres. Morbidity, including delayed gastric emptying, anastomotic leak and pancreatic fistula, remains high, affecting around 20–40 per cent of patients. Postoperative haemorrhage is less common, but is a life-threatening event with an estimated mortality rate of 20–50 per cent. Early-onset haemorrhage is rare, and generally occurs within 24 h, usually due to technical failures. Delayed haemorrhage occurring days or weeks after surgery occurs for a variety of reasons, but one cause of massive haemorrhage is from the formation of visceral arterial pseudoaneurysms. Although several arteries have been shown to be vulnerable to pseudoaneurysm formation, observational studies indicate that pseudoaneurysms of the gastroduodenal artery (GDA) are the most common.

The aim of this systematic review was to synthesize existing evidence regarding the incidence, clinical presentation and management of GDA pseudoaneurysms after pancreatic surgery.
Methods

Studies evaluating GDA pseudoaneurysm formation after pancreatic surgery were identified by means of database searches of MEDLINE and Embase. In Embase the terms used were: ‘false-aneurysm’ AND ‘gastroduodenal artery’ AND ‘pancreaticoduodenectomy’ OR ‘pancreatectomy’ OR ‘distal pancreatectomy’ OR ‘pylorus preserving pancreaticoduodenectomy’ OR ‘pancreas surgery’. In MEDLINE the terms used were: ‘false aneurysm’ OR ‘pseudoaneurysm’ AND ‘gastroduodenal artery’ AND ‘pancreaticoduodenectomy’ OR ‘pancreas surgery’ OR ‘pancreatectomy’ OR ‘distal pancreatectomy’ OR ‘pylorus preserving pancreaticoduodenectomy’.

A manual reference search was also performed to identify additional observational studies. No language restrictions were applied. Inclusion criteria were: manuscript published in a peer-reviewed journal until 2017, investigating adult patients aged over 18 years, undergoing pancreatic surgery for any indication, developing GDA pseudoaneurysm, and reporting clinical outcomes of interest. The authors independently reviewed all relevant titles and abstracts, and all disagreements were resolved by consensus. Observational studies that reported both the number of GDA pseudoaneurysms and the total number of pancreatic operations performed were used for quantitative analysis of incidence. Qualitative information also relevant to the clinical presentation and management of GDA pseudoaneurysms was extracted from individual cases and collated. Data extracted included index surgery, sentinel bleeding defined as haemorrhage that occurred in the gastrointestinal tract (intraluminal) or intra-abdominally (through a surgical drain) between 6h and 10 days before a massive haemorrhage in the postoperative setting, day of postoperative bleeding, diagnostic method, management,
Table 1 Studies included for determination of incidence

| Reference         | Country       | Study interval                  | Surgical procedures | Total no. of operations | Total no. of GDA pseudoaneurysms | Incidence (%) |
|-------------------|---------------|--------------------------------|---------------------|-------------------------|----------------------------------|---------------|
| Adam et al.       | Turkey        | January 1995 to January 2013   | PD                  | 342                     | 7                                | 2.0           |
| Suzuki et al.     | Japan         | January 2012 to July 2016      | PD                  | 88                      | 5                                | 6             |
| Jeong et al.      | South Korea   | October 1994 to December 2012  | PD                  | 1905                    | 18                               | 0.9           |
| Yada et al.       | Japan         | 1982–2010                      | PD + PPPD           | 361                     | 1                                | 0.3           |
| Loveček et al.    | Czech Republic| 2006–2015                      | PD                  | 449                     | 1                                | 0.2           |
| Fujii et al.      | Japan         | January 1993 to December 2005  | PD + PPPD + DP + SR + HPD + TP | 351 | 3 | 0.9 |
| Rajarathinam et al| India         | January 1998 to December 2007  | PD                  | 458                     | 2                                | 0.4           |
| Hur et al.        | South Korea   | March 2003 to March 2008       | PD                  | 192                     | 16                               | 8.3           |
| Sato et al.       | Japan         | January 1992 to December 1997  | PD                  | 81                      | 2                                | 2             |

Total: 4227 operations, 55 GDA pseudoaneurysms, incidence 1.3%

GDA, gastroduodenal artery; PD, pancreatoduodenectomy; PPPD, pylorus-preserving pancreatoduodenectomy; DP, distal pancreatectomy; SR, segmental resection; HPD, pancreatoduodenectomy plus hepatic resection; TP, total pancreatectomy.

Table 2 Data extracted for 39 patients with gastroduodenal artery pseudoaneurysm

| Reference         | Patient no. | Age (years) | Sex | Surgery | Complication       | Sentinel bleed | POD of bleed | Diagnostic method | Management | Survival |
|-------------------|-------------|-------------|-----|---------|---------------------|----------------|--------------|-------------------|------------|----------|
| Adam et al.       | 1           | 43          | M   | PD      | None                | Yes            | 45           | Angiography       | Selective embolization | Yes      |
|                   | 2           | 68          | F   | PD      | Abscess             | Yes            | 10           | Angiography       | Selective embolization | Yes      |
|                   | 3           | 59          | M   | PD      | Abscess             | Yes            | 4            | Angiography       | TAE of CHA | Yes      |
|                   | 4           | 41          | M   | PD      | Abscess             | Yes            | 23           | Angiography       | TAE of CHA | Yes      |
|                   | 5           | 63          | M   | PD      | Abscess             | Yes            | 14           | Angiography       | Selective embolization | No       |
|                   | 6           | 72          | F   | PD      | Abscess             | Yes            | 25           | Angiography       | Selective embolization | Yes      |
|                   | 7           | 51          | M   | PD      | None                | Yes            | 7            | Angiography       | TAE of CHA | Yes      |
| Fujii et al.      | 1           | –           | –   | HDP     | Pancreatic leak     | –              | 10           | Angiography       | TAE of CHA | Yes      |
|                   | 2           | –           | –   | PD      | Pancreatic fistula  | –              | 11           | Angiography       | Relaparotomy | Yes      |
|                   | 3           | –           | –   | HPD     | Pancreatic leak     | –              | 7            | Angiography       | Relaparotomy | No       |
| Rajarathinam et al| 1           | 52          | M   | PD      | Pancreatic fistula  | No             | 17           | Angiography       | Relaparotomy | Yes      |
|                   | 2           | 67          | M   | PD      | Intra-abdominal      | Yes            | 17           | Angiography       | Relaparotomy | No       |
| Hur et al.        | 1           | –           | –   | PPPD    | –                   | –              | 8            | –                 | TAE of CHA | No       |
|                   | 2           | –           | –   | PPPD    | –                   | –              | 6            | –                 | TAE of CHA | Yes      |
|                   | 3           | –           | –   | PD      | –                   | –              | 23           | –                 | Selective embolization | Yes      |
|                   | 4           | –           | –   | LPD     | –                   | –              | 15           | –                 | TAE of CHA | Yes      |
|                   | 5           | –           | –   | LPD     | –                   | –              | 7            | –                 | Selective embolization | Yes      |
|                   | 6           | –           | –   | PPPD    | –                   | –              | 12           | –                 | TAE of CHA | Yes      |
|                   | 7           | –           | –   | PPPD    | –                   | –              | 11           | –                 | Selective embolization | No       |
|                   | 8           | –           | –   | PPPD    | –                   | –              | 7            | –                 | TAE of CHA | Yes      |
|                   | 9           | –           | –   | PPPD    | –                   | –              | 8            | –                 | TAE of CHA | Yes      |
|                   | 10          | –           | –   | PPPD    | –                   | –              | 19           | –                 | TAE of CHA | Yes      |
|                   | 11          | –           | –   | PPPD    | –                   | –              | 7            | –                 | TAE of CHA | Yes      |
|                   | 12          | –           | –   | PPPD    | –                   | –              | 19           | –                 | TAE of CHA | Yes      |
|                   | 13          | –           | –   | PPPD    | –                   | –              | 8            | –                 | TAE of CHA | Yes      |
|                   | 14          | –           | –   | PPPD    | –                   | –              | 14           | –                 | TAE of CHA | Yes      |
|                   | 15          | –           | –   | HPD     | –                   | –              | 9            | –                 | TAE of CHA | Yes      |
|                   | 16          | –           | –   | PPPD    | –                   | –              | 13           | –                 | TAE of CHA | Yes      |
other postoperative complications and mortality. GDA pseudoaneurysms were confirmed either radiologically or during surgery in all studies.

Results

A PRISMA flow diagram is shown in Fig. 1. Some 88 studies were initially identified, 80 were screened, and 29 fulfilled the inclusion criteria. Of these, 13 studies\(^{12–24}\) were not included in the quantification as they were case series dealing exclusively with GDA pseudoaneurysms.

Five further studies were excluded as they did not provide the rate of GDA pseudoaneurysm in the postoperative period\(^{25–27}\), or the total number of pancreatic operations performed\(^{28,29}\). One study\(^{10}\) was excluded because it also included procedures not involving the pancreas (hepatic resection and gastrojejunostomy). Finally, one study\(^{9}\) was removed as it recruited patients from the same institution as another report\(^{31}\), but over a shorter period.

All of the nine manuscripts\(^{10,11,31–37}\) selected for quantitative analysis were single-centre observational studies (Table 1), mostly reporting on GDA pseudoaneurysms following pancreatoduodenectomy. One study\(^{13}\) included both pancreatoduodenectomy and modified pylorus-preserving pancreatoduodenectomy, while another\(^{10}\) included several pancreatic procedures. Data from 39 patients with GDA pseudoaneurysms were extracted from 14 studies for systematic review of clinical presentation and management (Table 2).

A total of 55 GDA pseudoaneurysms were identified in the postoperative period following 4227 pancreatic procedures, with a reported incidence of 1.3 (range 0.2–8.3) per cent (Table 1). Most patients who developed GDA pseudoaneurysm had a preceding complication in the postoperative period (14 of 21), including abscesses (6 patients), pancreatic fistulas (3) and pancreatic leaks (3). Three studies\(^{10,22,38}\) reported the formation of pseudoaneurysms away from the cut edge of the pancreas and in the absence of pancreatic fistulas. Sentinel bleeding was reported in 14 of 20 patients (70 per cent). The median time for postoperative haemorrhage was at 15 (range 4–210) days.

Diagnostic procedures were reported for 21 patients; 18 GDA pseudoaneurysms were detected by angiography.

| Reference | Patient no. | Age (years) | Sex | Surgery | Complication | Sentinel bleed | POD of bleed | Diagnostic method | Management | Survival |
|-----------|-------------|-------------|-----|---------|--------------|---------------|--------------|------------------|------------|----------|
| Miyazawa et al.\(^{13}\) | 1 | 71 | M | PD | Postoperative bleed | No | 180 | Contrast CT | Stenting | Yes |
| Lovećek et al.\(^{14}\) | 1 | 58 | M | PD | None | Yes | 18 | Angiography | Stenting | Yes |
| Mazza et al.\(^{16}\) | 1 | 61 | M | MSR | None | No | 210 | Contrast CT | Selective embolization | Yes |
| Huang et al.\(^{17}\) | 1 | 72 | M | PD | – | Yes | 17 | – | Selective embolization | Yes |
| Huang et al.\(^{17}\) | 2 | 65 | F | Duodenum-preserving pancreatic resection | – | Yes | 30 | – | Selective embolization | No |
| Noun et al.\(^{18}\) | 1 | 58 | M | PD | Pancreatic fistula | Yes | 19 | Angiography | Selective embolization | Yes |
| Orsenigo et al.\(^{21}\) | 1 | 38 | M | SPK | AV fistula | No | 15 | MR angiography | Selective embolization | Yes |
| Sugimoto et al.\(^{22}\) | 1 | 62 | M | PD | None | No | 120 | Angiography | TAE of CHA | Yes |
| Born et al.\(^{23}\) | 1 | 42 | M | Lateral pancreateo-jejunostomy | None | Yes | 21 | Angiography | TAE of CHA | Yes |
| Teramoto et al.\(^{24}\) | 1 | 70 | M | PD | Pancreatic leak | No | 34 | Angiography | Selective embolization | Yes |
| Okuno et al.\(^{30}\) | 1 | 46 | F | PD | None | Yes | 62 | Angiography | Selective embolization | Yes |
| Overall | 39 | 60* | 16 M, 4 F | | | 14 of 21 | 14 of 20 | 15 (4–210)* | 33 of 39 |

*Median (range) value. POD, postoperative day; PD, pancreatoduodenectomy; TAE, transarterial embolization; CHA, common hepatic artery; HPD, pancreatectomy plus hepatic resection; PPPD, pylorus-preserving pancreatoduodenectomy; LPD, laparoscopic pancreatoduodenectomy; MSR, middle segment resection; SPK, simultaneous pancreas–kidney transplant.
Thirty-five of the 39 patients (90 per cent) were treated using an endovascular approach. Nineteen (49 per cent) were managed using transarterial embolization (TAE) of the GDA via the common hepatic artery (CHA), and 14 (36 per cent) by selective embolization of the pseudoaneurysm. Stenting was employed in two patients (5 per cent), and only four (10 per cent) were treated by emergency laparotomy. The overall survival rate was 85 per cent (33 of 39).

Discussion

The GDA is the most common site for pseudoaneurysm formation after pancreatic surgery9–11, and its rupture in the postoperative period has long been recognized as a cause of substantial morbidity and mortality2,8. GDA pseudoaneurysms are rare. The present analysis suggests that they occur in 0·2–8·3 per cent of pancreatic resections. It should be noted, however, that studies included in this review were all high-volume resectional centres.

In this series, two-thirds of the patients (4 of 21) had a preceding complication following pancreatic resection. Most authors favoured the hypothesis that lytic, enzyme-rich, pancreatic fluid from a pancreatic anastomotic leak could result in autodigestion of GDA vessel wall owing to its proximity to the pancreatic anastomosis. Interestingly, a few studies10,22,38 reported the formation of pseudoaneurysms at distance from the pancreatic anastomosis and in the absence of an overt pancreatic fistula, suggesting that minor iatrogenic injury, such as skeletonization of the vessel wall during extensive lymphadenectomy, may lead to vessel weakening and subsequent pseudoaneurysm formation.

Various techniques have been suggested to reduce the chance of pseudoaneurysm formation, including the ‘wrapping’ technique25,39–45, in which the exposed retroperitoneal vessels are covered with omentum or the falciform ligament. Others46 have suggested leaving 1 cm at the origin of the GDA stump to minimize the likelihood of lytic pancreatic juices coming into contact with the vessel.

Recognition of a sentinel bleed may help in early management, this being a feature in most patients9,37,47. Although sentinel bleeding was associated with poor outcome in some series6,35, few authors discussed the importance of immediate angiography after a sentinel bleed to look for the possibility of a ruptured pseudoaneurysm6,37. Although angiography also has the added benefit of allowing transition to endovascular treatment, a number of reports noted that negative findings cannot be used to exclude a bleeding pseudoaneurysm26, as bleeding can be intermittent or the rate of bleeding is below the detection limit of the equipment7,9,37,48,49.

Surgical intervention has been largely replaced by interventional radiology50–54. Some older studies26,55 advocated surgery in the context of additional intra-abdominal complications such as pancreatic fistula, but more recent series11,36 have documented the superiority of endovascular management. A recent meta-analysis46 of non-randomized studies comparing endovascular management and laparotomy for delayed massive haemorrhage suggested lower complication and mortality rates in the endovascular group.

The endovascular management of pseudoaneurysms varied, reflecting the location and size of the pseudoaneurysm and probably institutional preferences for approach and embolization technique, and materials. TAE of the GDA was conducted via either the CHA or the superior mesenteric artery to achieve both proximal and distal occlusion, to exclude the pseudoaneurysm and prevent backflow from collateral circulation57. Such an approach should consider patency of the portal venous system51 as TAE distal and proximal to the GDA pseudoaneurysm can cause complete occlusion of the CHA, leading to liver infarction (reported range 30–66 per cent)29,37,58, as well as hepatic failure and abscess formation58. Covered stents represented the alternative to TAE. The key advantage over TAE would be in maintaining patency of the CHA and reducing the risk of hepatic infarction, although accurate stent deployment might be technically more challenging and time-consuming than TAE36,57. Despite these issues, stenting seems to be preferred in more recent series, on the grounds that selective embolization of the GDA stump or pseudoaneurysm seems to be associated with high rates of recurrent bleeding36,42.

Limitations of this study include heterogeneity of the included studies, the descriptors used and study sizes. The absence of any prospective registers or clinical trials on this topic needs to be addressed.

Disclosure

The authors declare no conflict of interest.

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