Para-anastomotic aneurysm (PAAA) of the abdominal aorta is a major complication of infrarenal grafting for occlusive disease or aneurysm, and involves the suture-line with disruption of the anastomosis. PAAA may be related to infection of the graft or caused by a sterile degenerative process involving the anastomosis. It is rare for a PAAA to be diagnosed within 5 years after aortic grafting (1-6%), and such cases are generally associated with an infection of the graft; the uninfected degenerative form of PAAA is usually observed 8-10 years after primary aortic surgery.

We focused our attention on uninfected PAAA because such aneurysms are secondary to a slow, degenerative process acting silently at the anastomosis, apparently not influenced by any external factors, and responsible for late failure of aortic reconstructions with life-threatening clinical consequences. A PAAA may present as a true aneurysm of the aortic remnant near the anastomosis, or as a false aneurysm (anastomotic aneurysm) of the suture-line, and both are potentially responsible for thrombosis, distal embolization, compression/erosion of adjacent organs, and inexorable growth to the point of rupture.

Their deep location makes PAAA difficult to detect without imaging and, despite the

**Purpose:** This single-institution retrospective review examines the management of uninfected para-anastomotic aneurysms of the abdominal aorta (PAAA), developed after infrarenal grafting. **Materials and Methods:** From October 1979 to November 2005, 31 PAAA were observed in our Department. Twenty-six uninfected PAAA of degenerative etiology, including 24 false and 2 true aneurysms, were candidates for intervention and retrospectively included in our database for management and outcome evaluation. Six (23%) patients were treated as emergencies. Surgery included tube graft interposition (n = 12), new reconstruction (n = 8), and graft removal with extra-anatomic bypass (n = 3). Endovascular management (n = 3) consisted of free-flow tube endografts. **Results:** The mortality rate among the elective and emergency cases was 5% and 66.6%, respectively (p = 0.005). The morbidity rate in elective cases was 57.8%, whereas 75% in emergency cases (p = 0.99). The survival rate during the follow-up was significantly higher for elective cases than for emergency cases. **Conclusion:** Uninfected PAAA is a late complication of aortic grafting, tends to evolve silently and is difficult to diagnose. The prevalence is underestimated and increases with time since surgery. The mortality rate is higher among patients treated as an emergency than among patients who undergo elective surgery, therefore, elective treatment and aggressive management in the case of pseudoaneurysm are the keys to obtain a good outcome. Endovascular treatment could reduce mortality. Patients who undergo infrarenal aortic grafting require life-long surveillance after surgery.

**Key Words:** Abdominal aortic grafting, pseudoaneurysm, anastomotic aneurysm, para-anastomotic aneurysm, endograft
high levels of sensitivity and specificity of diagnostic techniques now available, true and false PAAA often cannot clearly be distinguished. Furthermore, the follow-ups scheduled by most Departments of Vascular Surgery to check aortic grafts are not continued for long enough period, considering the time at which uninfected PAAA usually develops. For these reasons, the real frequency of uninfected PAAA is difficult to determine. Based on the limited data available in the literature, the overall incidence appears to range from 0.2% to 15%, however, this is probably an underestimate because of no long-term, standardized follow-ups that could reveal a higher frequency than that so far reported. One further problem is that many of the published epidemiological data have been derived from heterogeneous populations of patients with no distinctions concerning the etiology or site of the disrupted anastomoses.

Prospective analyses estimated that the incidence of late, uninfected anastomotic complications ranges from 22.8% to 35.8% at 15 years after infrarenal grafting. Such results can not certainly be negligible, and raise doubts about the long-term durability of what has always been considered the most successful and gratifying intervention by vascular surgeons.

Once diagnosed, the management of uninfected PAAA is extremely challenging both in elective and emergency conditions, and the outcome appears to be influenced more by technical surgical variables than by the patients’ clinical background. The high morbidity and mortality rates reported in the literature, particularly following surgery performed as an emergency, bear witness to the clinical relevance of this complication and the importance of an early diagnosis in order to manage it in elective conditions.

The aim of this single-institution study was to review the management of uninfected PAAA in our Department, focusing on perioperative outcomes and long-term survival.

**MATERIALS AND METHODS**

All the infrarenal aortic prosthetic reconstructions performed in our Institution between October 1979 and November 2006 were identified from the prospective patients’ registry, and a retrospective review of patients’ clinical records was used to create a database containing information on all the cases of PAAA observed, including anatomical and etiopathogenetic details of the PAAA based on the results of clinical and imaging evaluations. Infected PAAA were excluded.

Pre-operative medical risk factors, intra-operative variables, and post-operative outcomes were analyzed for uninfected degenerative PAAA only. Pre-operative risk factors were recorded according to the American Society of Anesthesiologists (ASA) physical status classification after evaluation of cardiopulmonary and renal function. These data together with indications for the operation and type of treatment allowed to calculate the risk/benefit ratio and identification of patients unfit for an intervention. All complicated, life-threatening PAAA, were treated as emergencies without evaluation of pre-operative risk factors. Operative variables recorded included the form (elective or emergency) and type (surgical or endovascular) of the intervention, type of surgical reconstruction, and anatomical location of the aortic control. Post-operative outcomes included in-hospital morbidity and mortality: in-hospital mortality was defined as a peri-operative death occurring during the same admission regardless of the duration of the hospital stay; in-hospital morbidity included cardiac, pulmonary, neurological, gastrointestinal, visceral ischemic, peripheral ischemic, systemic, and wound complications occurring during the same hospital admission. Long-term survival of more recent cases was assessed by direct clinical evaluation, and by telephone contact for older ones.

The primary end-point for analysis was perioperative outcome, and the secondary end-point was long-term survival. Demographic data, pre-operative medical risk factors, and peri-operative variables were analyzed descriptively. Mortality and morbidity rates were considered separately for the overall population of patients with uninfected PAAA, and the subgroups were treated electively or as an emergency and those managed surgically or with an endovascular intervention. Fisher’s exact test was used to compare the results of patients treated electively or as an emergency and those undergoing surgery or endovascular treatment (two-tailed level of significance $\alpha = 0.05$). The patients who were managed surgically underwent physical examination after 2 months and subsequently once yearly; data from these examinations were combined with those from ultrasonography when diagnostic echo-studies became widely available in clinical practice; angio-computed tomography (angio-CT) was performed only if abnormalities were suspected at ultrasonography and routinely after 5 years. Patients managed with an endovascular procedure underwent angio-CT after 2, 6, and 12 months and then yearly. The patients’ survival was assessed using Kaplan-Meier product-limit estimates (SPSS 13.0, Chicago, IL, USA). $p$ values < 0.05 were considered statistically significant.

**RESULTS**

From October 1979 to November 2005, 2,996 infrarenal prosthetic reconstructions for aneurysmal and obstructive aortoiliac disease were performed in our Department. The demographics of the population who underwent these reconstructions are presented in Table 1.

During the same period, 31 patients with PAAA were observed. Seventeen patients (55%) had had their primary graft in our Department, whereas 14 patients (45%) were treated at another Institution. All patients underwent clinical, biochemical, and microbiological examinations along with imaging studies in order to obtain a precise diagnosis and define the
etiology and morphology of the PAAA. The type of imaging study (thoraco-abdominal angio-CT, abdominal color Doppler ultrasound, aortography) varied over time because of technological evolutions that occurred during the 26-year period covered in this study.

Four of the 31 patients with an identified PAAA had graft infection combined with disruption of the proximal aortic anastomosis and formation of a pseudoaneurysm. The remaining 27 patients had uninfected PAAA, as determined by clinical evaluation, laboratory examinations, microbiological analyses and imaging, thereby excluding an infective etiology, and anastomotic failure was the result of a degenerative process.

Table 1. Demographics of Population Underwent Infrarenal Aortic Grafting

| Aneurysm | Obstructive disease |
|----------|---------------------|
| Age (yrs) | 71.2 ± 7.6 | 64.3 ± 9.2 | 67.5 ± 5.1 |
| N° | % | N° | % |
| N° | 2,486 | 83 | 510 | 17 | 2,996 |
| Male | 2,337 | 94 | 389 | 76 | 2,726 (91%) |
| Hypertension | 2,287 | 92 | 449 | 88 | 2,736 (91%) |
| Dislipidemia | 1,516 | 61 | 342 | 67 | 1,858 (62%) |
| Smoking history | 1,989 | 80 | 449 | 88 | 2,438 (81%) |
| Diabetes | 597 | 24 | 133 | 26 | 730 (24%) |
| Cardiac disease | 1,616 | 65 | 270 | 53 | 1,886 (63%) |
| CAD | 1,342 | 54 | 250 | 49 | 1,592 (53%) |
| COPD | 820 | 33 | 132 | 26 | 952 (32%) |
| Cerebrovascular disease | 472 | 19 | 66 | 13 | 538 (18%) |
| CRF | 597 | 24 | 117 | 23 | 714 (24%) |
| Peripheral vascular disease | 547 | 22 | 270 | 53 | 817 (27%) |

COPD, chronic obstructive pulmonary disease; CAD, coronary artery disease; CRF, chronic renal failure.

Table 2. Comorbidities and Risk Factors

| N° | % | PAAA | Complicated cases (n = 10) |
|----|---|------|---------------------------|
| Smoking history | 13 | 46.4 | 12 false 1 true | 4 | 40 |
| COPD | 14 | 50 | 14 false | 5 | 50 |
| Hypertension | 11 | 39.2 | 10 false 1 true | 4 | 40 |
| CAD | 6 | 21.4 | 6 false | 4 | 40 |
| Atrial fibrillation | 7 | 25 | 7 false | 3 | 30 |
| Left ventricular hypertrophy | 2 | 7.14 | 2 false | 2 | 20 |
| Pace-maker | 2 | 7.14 | 2 false | 1 | 10 |
| Dilatative cardiomyopathy | 3 | 10.7 | 3 false | - | - |
| CRF | 7 | 25 | 7 false | 3 | 30 |
| Lower limb obstructive disease | 8 | 28.6 | 6 false 2 true | 3 | 30 |
| Stroke | 2 | 7.14 | 2 false | - | - |
| Dyslipidemia | 5 | 17.8 | 3 false 2 true | - | - |
| Diabetes | 2 | 7.14 | 2 false | - | - |
| Other false aneurysms | 7 | 25 | 7 false | 1 | 10 |
| Other true aneurysms | 3 | 10.7 | 3 false | - | - |
| Oral anticoagulants | 4 | 14.3 | 4 false | 2 | 20 |

PAAA, para-anastomotic aneurysm of the abdominal aorta; COPD, chronic obstructive pulmonary disease; CAD, coronary artery disease; CRF, chronic renal failure.
All patients in whom an uninfected PAAA was detected in non-emergency conditions underwent preoperative assessment of cardiac, pulmonary, renal function (color Doppler echocardiography, coronarography, spirometry, and serum creatinine levels), and comorbid conditions in order to stratify the operative risk according to the ASA criteria. For patients presenting as an emergency, there was usually not enough time for a pre-operative assessment, except for that strictly related to the diagnosis of PAAA. Emergency cases underwent immediate treatment, whereas for patients whose PAAA was detected in non-emergency conditions the indication to treat was discussed considering the ASA score, the diameter of the PAAA and the risk related to technical-surgical variables, in order to define a benefit/risk ratio.

Our database of patients with uninfected PAAA included information on 25 (92.6%) men and 2 women. The mean age of these patients was 70 years (range, 47 to 87 years), whereas the mean age at the time of primary surgery was 58 years. The mean interval from primary surgery to the diagnosis of PAAA was 10.1 years (range, 0.25-21 years).

The indication for primary grafting was aneurysm in 18 patients and occlusive disease in 9 patients. The patients’ comorbidities and risk factors are listed in Table 2.

After careful preoperative evaluation, 25 patients - with 26 uninfected PAAA - underwent intervention: 24 (92.3%) with pseudoaneurysms - 23 proximal, 1 distal - and 2 with true aneurysms. Two proximal false PAAAs were associated with Crawford’s type 4 thoraco-abdominal aortic aneurysm (TAA), and no other thoracic aortic aneurysm recurrence was diagnosed in the population treated. In one patient, a false PAAA recurred twice at the same anastomotic line. Two asymptomatic patients were considered unfit for intervention: 1 was an 83-year-old man with a proximal pseudoaneurysm and type 3 TAA 90 mm in diameter, who had a ventriculoperitoneal drainage for a hydrocele. He was permanently confined to bed because of a spinal cord disease, and had severe chronic obstructive pulmonary disease and cardiac dysfunction. The other patient was an 87-year-old man with a proximal pseudoaneurysm of 42 mm in diameter who had severe chronic obstructive pulmonary disease, mild-to-severe renal impairment and extreme cardiac dysfunction (ejection fraction 18% at echocardiography).

Among the patients with a false PAAA, the disrupted anastomosis was end-to-side in 9 cases and end-to-end in the other 15 (62.5%) cases. In the 2 patients with true PAAA, the disrupted anastomosis was end-to-end in one and end-to-side in the other. The mean diameter of the true PAAA was 50 mm (range, 48-52 mm) whereas that of the false PAAA was 46.25 mm (range, 34-80 mm). The mean diameter of the PAAA treated as emergencies because of rupture or aorto-enteric fistula (AEF) was 45 mm (range, 34-53 mm).

On admission to the hospital, patients with true PAAA were asymptomatic and free of complications. Thirteen (50%) of the patients with false PAAA had acute or subacute non-specific symptoms, including abdominal pain, gastrointestinal bleeding, lower limb ischemia, asthma and back pain with sciatica.

In the subgroup of false PAAA, 10 (41.6%) cases were complicated by thrombosis of the aneurysm and lower limb ischemia (n = 1), asymptomatic fissuration diagnosed intraoperatively (n = 2), asymptomatic tamponed rupture in mesocolon (n = 1), retroperitoneal rupture (n = 2) and secondary AEF with upper gastrointestinal bleeding (n = 4). True secondary AEF in the third part of the duodenum was observed in 2 cases of ruptured PAAA, whereas a paraprosthetic secondary AEF with duodeno-ileal perforation over an integral pulsating PAAA capsule was detected in another 2 cases.

Clinical examination was negative in 9 cases (35%), whereas a pulsating abdominal mass was palpable in 17 cases (65%).

Ultrascanography was diagnostic in 4 of the 9 false PAAA in which this examination was used and the aortic dilatation was detected without defining its extension. Angio-CT was diagnostic in all the cases in which it was applied and the extent of the PAAA and the involvement of visceral arteries were determined. Aortography was diagnostic in 16 (80%) of 20 cases (19 pseudoaneurysms and 1 true aneurysm). Esophagogastroduodenoscopy was performed in 3 cases of hematemesis of duodenal origin, but failed to detect fistulae. One false PAAA was diagnosed during an explorative laparotomy.

In the subgroup of patients treated surgically, no infective foci were found during intra-operative exploration. Samples of the grafts removed were microbiologically and histologically examined, which confirmed the lack of bacterial contamination and demonstrated the degenerative etiopathogenesis.

AEF is potentially responsible for suture-line infection from the bowel. The absence of infective features might be explained by the fact that the true AEF after PAAA rupture (n = 2) developed so acutely that there was no time for contamination of the anastomosis to occur, whereas the integrity of the false PAAA fibrous capsule probably protected the aortic suture-line from infection in paraprosthetic AEF (n = 2). Furthermore, the bacterial load of the duodenum is low compared to that of other parts of the intestine.

Twenty cases (77%) - 2 true and 18 false PAAA - were treated electively, whereas 6 symptomatic and life-threatening complicated false PAAA were treated in emergency. Table 3 summarizes the presentation and management of the cases of PAAA.

Surgery was performed in 23 cases (88%). The transabdominal approach allowed good exposure of the entire abdominal aorta in 21 cases (20 false and 1 true PAAA). Aortic control was safely obtained below or close to renal arteries, except in 6 cases which required suprarenal clamping below the mesenteric vessels because of difficulties in the surgical approach to the juxtarenal aorta and PAAA. The thoracoabdominal approach was required for 2 false PAAA associated with type 4 TAA.
Ten cases of false PAAA were treated by in situ Dacron reconstruction with tube graft interposition between the aorta and the distal portion of the old graft; 7 cases of false PAAA and 1 true PAAA were managed by complete removal of the old graft and new reconstruction; thoraco-abdominal endoaneuysmectomy and tube graft interposition between the thoracic aorta and old graft with reimplantation of visceral arteries was performed in two cases of false PAAA with type 4 TAA. Extra-anatomic axillofemoral bypass with reinforced expanded PolyTetraFluorEthilene (ePTFE) combined with complete graft removal and closure of the aortic stump was the management chosen for 3 cases of PAAA, in order to avoid as much as possible technical risks related to in situ reconstruction in the site of a damaged, malacic aorta or when there was a suspicion of bacterial contamination from the duodenal lumen at the onset of AEF.

Endovascular treatment was performed electively in 3 uncomplicated cases (2 proximal false PAAA and 1 proximal true PAAA after end-to-end anastomosis) in the operating room, after surgical exposure of the femoral arteries under

Table 3. Presentation and Management of Un-Infected PAAA

| PAAA     | Ø (mm) | Clinics             | Complications      | Treatment            | In-hospital outcome |
|----------|--------|---------------------|--------------------|----------------------|---------------------|
| 1 False  | 45     | Asymptomatic        | -                  | Elective             | Tube graft interposition | Alive           |
| 2 False  | 53     | Abdominal pain, shock | Retroperitoneal rupture | Emergency            | Tube graft interposition | Dead            |
| 3 False  | 50     | Lower limb ischemia | -                  | Elective             | New reconstruction    | Alive           |
| 4 False  | 40     | Asymptomatic        | -                  | Elective             | New reconstruction    | Alive           |
| 5 False  | 52     | Abdominal pain      | Fissurisation*     | Elective             | Tube graft interposition | Alive           |
| 6 False  | 50     | Abdominal pain, shock | Retroperitoneal rupture | Emergency            | Graft removal and extraanatomic bypass | Dead            |
| 7 False  | 40     | Asymptomatic        | -                  | Elective             | Tube graft interposition | Alive           |
| 8 False  | 40     | Abdominal pain      | Fissurisation*     | Elective             | New reconstruction    | Alive           |
| 9 False  | 38     | Asymptomatic        | -                  | Elective             | Graft removal and extraanatomic bypass | Alive           |
| 10 False | 75     | Asymptomatic        | -                  | Elective             | Tube graft interposition | Alive           |
| 11 False | 40     | Asymptomatic        | -                  | Elective             | Tube graft interposition | Alive           |
| 12 False | 50     | GI bleeding, shock  | True AEF           | Emergency            | Tube graft interposition | Alive           |
| 13 False | 42     | Asymptomatic        | -                  |                     | Unfit for intervention |                |
| 14 True  | 52     | Asymptomatic        | -                  | Elective             | New reconstruction    | Alive           |
| 15 False | 34     | GI bleeding         | Paraprosthetic AEF | Emergency            | Tube graft interposition | Alive           |
| 16 False | 38     | Asymptomatic        | -                  | Elective             | New reconstruction    | Alive           |
| 17 False | 47     | GI bleeding         | Paraprosthetic AEF | Emergency            | New reconstruction    | Dead            |
| 18 False | 41     | Abdominal pain      | -                  | Elective             | Tube graft interposition | Alive           |
| 19 False | 43     | Asymptomatic        | Tamponated rupture in mesocolon | Elective | New reconstruction | Alive           |
| 20 False | 36     | Asymptomatic        | -                  | Elective             | Tube graft interposition | Alive           |
| 21 False | 36     | GI bleeding, shock  | True AEF           | Emergency            | Graft removal and extraanatomic bypass | Dead            |
| 22 False | 37     | Lower limb ischemia | Thrombosis         | Elective             | New reconstruction    | Alive           |
| 23 False | 50     | Lumboischialgia      | -                  | Elective             | Endograft             | Alive           |
| 24 True  | 48     | Asymptomatic        | -                  | Elective             | Endograft             | Alive           |
| 25 False | 35     | Asymptomatic        | -                  | Elective             | Endograft             | Alive           |
| 26 False and TAA | 80 | Abdominal pain | -                  | Elective             | Tube graft interposition | Alive           |
| 27 False and TAA | 60 | Asymptomatic | -                  | Elective             | Tube graft interposition | Dead            |
| 28 False and TAA | 90 | Asymptomatic | -                  |                      | Unfit for intervention |                |

PAAA, para-anastomotic aneurysm of the abdominal aorta; GI, gastrointestinal; TAA, thoracoabdominal aortic aneurysm.

*Intraoperative diagnosis.
epidural anesthesia. Angio-CT scanning and complementary aortography documented proximal infrarenal necks shorter than 15 mm, so free-flow tube endografts (Endologix Powerlink Bard®) were required to achieve safe exclusion of the aneurysm without impairing renal vessels.

Five patients, all treated surgically, died in hospital giving an in-hospital mortality rate of 19.2% in the whole series and 21.7% in the surgical subgroup. No deaths occurred in the endovascular subgroup. The mortality rate was 5% among patients treated electively and 66.6% in emergency cases (\(p = 0.005\), 2-tailed Fisher’s exact test). The mortality rate among patients who underwent localized and extensive surgical interventions was 0% and 50%, respectively (\(p = 0.09\), two-tailed Fisher’s exact test). The mortality rate in cases of symptomatic PAAA was 30.7%, whereas it was 7.7% in asymptomatic cases (\(p = 0.32\), two-tailed Fisher’s exact test). One patient with a false PAAA and type 4 TAA died during an elective intervention from consumption coagulopathy after severe aortic bleeding. Four patients treated as emergencies died in the post-operative period: 2 from compromised hemodynamics and coagulopathy the first day after surgery for ruptured false PAAA; 1 patient died from massive bowel infarction 8 days after surgery for false PAAA and secondary AEF; and 1 died from acute massive duodenal bleeding and perforation 29 days after surgery for false PAAA complicated by gastrointestinal bleeding.

The overall morbidity rate was 66.5%; 57.8% in electively managed cases and 75% in emergency cases (\(p = 0.99\), 2-tailed Fisher’s exact test). The morbidity rate in symptomatic cases was 63.6%, whereas it was 50% in the asymptomatic group (\(p = 0.68\), two-tailed Fisher’s exact test). Major and minor complications observed are summarized in Table 4. Only minor complications were noted in the endovascular subgroup, and their relationship with the procedure appeared very weak.

Technical difficulties were recorded in 21 cases of redo-surgery: omental and bowel adhesions to PAAA with or without AEF (\(n = 12\)); challenging preparation of the aorta, PAAA and renal arteries due to intense fibrosis (\(n = 5\)); ligature of the left renal vein to gain the aortic clamp-site (\(n = 2\)); iatrogenic duodeno-ileal damage (\(n = 3\)); retroperitoneal/abdominal bleeding after PAAA rupture or consumption coagulopathy (\(n = 4\)); intra-operative rupture of the aneurysm during preparation of the aorta (\(n = 1\)).

A stay in the Intensive Care Unit (ICU) was required in 41.6% of cases for a mean period of 2.5 days (range, 1-8 days). None of the patients in the endovascular subgroup needed ICU management. Survivors were discharged from hospital after a mean period of 12.2 days (range, 4-27 days); 14.3 and 4.3 days

Table 4. Early Post-Operative Complications

| Events                                      | No | PAAA                      | Clinics                  | Treatment                | Outcome                      |
|---------------------------------------------|----|---------------------------|--------------------------|--------------------------|------------------------------|
| Cardiac arrest                              | 1  | Distal false ruptured and tamponated | Asymptomatic             | Elective surgery         | Recovery after resuscitation |
| Pneumothorax                                | 2  | Proximal false            | Asymptomatic             | Elective surgery         | Recovery                     |
| Massive intestinal infarction               | 1  | Proximal false and AEF    | Hematemesis and shock    | Emergency surgery        | Death                        |
| Duodenal ulcercation/perforation            | 1  | Proximal false and duodenal ulceration | Hematemesis and melena | Emergency surgery        | Death for massive bleeding   |
| Minor stroke (hypotension related)          | 1  | Proximal false ruptured in duodenum | Hematemesis              | Emergency surgery        | Recovery                     |
| Acute lower limb ischemia                   | 1  | True                      | Chronic lower limbs ischemia | Elective surgery         | Recovery after thrombectomy  |
| Laparotomy’s dehiscence and evisceration    | 1  | Proximal false with thrombosis | Subacute lower limbs ischemia | Elective surgery         | Recovery after laparoplasty  |
| Transient renal failure                     | 2  | Proximal false            | Asymptomatic             | Elective surgery         | Recovery                     |
| Severe hypotension                          | 2  | Proximal false ruptured in duodenum | Asymptomatic/ Hematemesis | Elective/Emergency surgery | Recovery                     |
| Severe anemia                               | 3  | Proximal false            | Asymptomatic             | Elective surgery/ EVG    | Recovery after transfusions  |
| Groin lymphorrhoea                          | 1  | Proximal false in fissurisation | Abdominal pain          | Elective surgery         | Recovery                     |
| Acute podagra                               | 1  | True                      | Asymptomatic             | EVG                      | Recovery                     |

PAAA, para-anastomotic aneurysm of the abdominal aorta; AEF, aorto-enteric fistula; EVG, endovascular grafting.
for the surgical and endovascular subgroups, respectively. Follow-up data are available for all the patients. Kaplan-Meier product-limit survival estimates are depicted in Figs. 1, 2 and 3. With a median follow-up of 105.6 months (range, 12-312 months), the 10-year survival was 50% for the overall population of patients with uninfected PAAA who underwent an intervention (Fig. 1). There was a significant difference in mid-to long-term survival between elective and emergency cases, as depicted in Fig. 2. Eight patients died of causes unrelated to the PAAA treatment. There were no cases of graft infection or late lower limb amputations. No graft complications were observed in the surgical subgroup except for that of the patient who experienced recurrent false PAAA. Three patients underwent interventions for disease of other vascular districts. One pseudoaneurysm treated with an endograft developed a proximal type I endoleak, detected by angio-CT 2 months after the procedure: an aortic cuff with effective exclusion of the aneurysm was deployed in this patient. Exclusion of the aneurysm was maintained in the other 2 cases in which endovascular management was used without signs of stent-graft migration or failed fixation to the inner surface of the old grafts. The 2 patients considered unfit for intervention died; one after rupture of a type 3 TAA and the other of cardiac failure and uremia.

**DISCUSSION**

Uninfected PAAA is a late major complication of infrarenal aortic grafting. Discontinuity of the anastomosis causes a false PAAA that usually occurs as a result of arterial wall disruption and mechanical stress rather than because of graft/suture material failure. Aorta versus graft compliance/diameter mismatch and extreme mechanical stress (greater in end-to-side rather than end-to-end anastomoses) at the suture-line play major pathogenetic roles. Hypertension, atherosclerosis, and previous aortic endarterectomy are important in suture-line failure. Major peri-operative complications, technical errors, recurrent aneurysm and α-antitrypsin deficiency may be associated with an early onset of pseudoaneurysm.

True PAAA usually occurs after surgery performed for an aneurysm rather than for occlusive disease, and is a progression of the aortic dilatation it may affect the infrarenal aorta as a consequence of inadequate resection of the primary aneurysm, particularly when treated as an emergency for rupture, or it involves the pararenal and/or suprarenal aorta proximally to a graft placed just below the renal arteries as a consequence of progressive atherosclerosis. The incidence of false PAAA is 2 to 3 times higher than that of true PAAA.

Uninfected PAAA is, however, an underestimated complication. Table 5 summarizes the reported incidences of PAAA in different series. There is a tendency to define PAAA as an infrequent event, but there is a clear lack of uniformity of
data, therefore, a precise epidemiological evaluation of the phenomenon is not possible.\textsuperscript{11} The real dimensions of this complication are probably misunderstood\textsuperscript{7,9,18} because of the heterogeneous attention paid to PAAA by Departments of Vascular Surgery, the lack of long-term follow-ups after aortic grafting, diagnostic difficulties (PAAA evolves silently for a long time and manifests with non-specific symptoms and no signs because of its retroperitoneal location), and the different diagnostic methods used during follow-up.\textsuperscript{3,6,7,9,10} The significant number of PAAA (45\% in our survey) observed in Institutions different from those in which the primary aortic reconstructions were performed confirms that it is difficult to achieve long-term follow-up after aortic grafting by many Vascular Departments.\textsuperscript{8,32} In our own experience, we diagnosed PAAA in 17 patients from a population of 2,996 who underwent infrarenal reconstructions, resulting in an incidence of 0.57\% PAAA. We consider this an underestimate, because we were not able to follow-up all the patients over the 26-year period and we cannot determine how many developed late anastomotic complications that were eventually diagnosed and treated at other Departments of Vascular Surgery.

Prospective analyses\textsuperscript{10,13,14} underline how the incidence of PAAA increases significantly with time elapsed since the primary graft, so that the patient’s long life-expectancy is a risk factor for future development of sterile PAAA.\textsuperscript{3,4} Using life-table analysis, Edwards et al.\textsuperscript{10} estimated that the overall incidence of PAAA after 8 and 15 years was 5\% and 27\%, respectively, and that the incidence of false PAAA was 1\% and 20\%, respectively. They also estimated that the incidence of true/recurrent PAAA was 4\% and 9\% after 10 and 25 years, respectively. Van den Akker et al.,\textsuperscript{13} who also used life-table analysis, calculated that the chance of being free of an anastomotic aneurysm at any site was 77.2\% at 15 years after the primary operation. By applying the Kaplan-Meier method, Mii

| Table 5. PAAA’s Incidence in Literature |
|----------------------------------------|
| Casistics | PAAA | False PAAA | True PAAA |
|-----------------|-----------------|-----------------|-----------------|
| Szilagyi\textsuperscript{19} | Global | - | 1.7\% per anastomotic site | - |
| | | | 0.2\% per aortic anastomosis | - |
| Starr\textsuperscript{21} | Global | - | 2\% per patient | - |
| Millili\textsuperscript{20} | AAA | - | 4.5\% per patient | - |
| | AOD | - | 4.5\% per patient/femoral site | - |
| | | | 0.7\% per patient/aortic site | - |
| Plate\textsuperscript{25} | AAA | 5.4\% per patient | 1.3\% per patient | 2.9\% per patient (abdominal and thoraco-abdominal) |
| Mehigan\textsuperscript{22} | Global | - | 2.7\% per anastomosis | - |
| | | | 1.5\% per anastomosis/AAA | - |
| | | | 5.2\% per anastomosis/PAOD | - |
| Szilagyi\textsuperscript{23} | AOD | - | 3.7\% per anastomosis | - |
| | | | 0.2\% per aortic anastomosis | - |
| Signorelli\textsuperscript{15} | Global | - | 2\% per anastomotic site | - |
| | | | 0.16\% per aortic anastomosis | - |
| Nevelsteen\textsuperscript{31} | AOD | - | 1.4\% per patient (retroperitoneal site) | - |
| van der Akker\textsuperscript{13} | AOD | - | 13.3\% per patient | - |
| | | | 4.8\% per aortic anastomosis | - |
| Mikati\textsuperscript{24} | AOD | - | 15\% per patient (aortic anastomosis) | - |
| Edwards\textsuperscript{20} | AAA | 10\% per patient (intraabdominal) | - | 8\% per patient (only after surgery for AAA) |
| | AOD | - | - | - |
| Melliere\textsuperscript{25} | AAA | - | 2-5\% | - |
| | AOD | - | 3\% per patient | 13\% per patient (supraanastomotic aorta) |
| | | | 1\% per patient/aortic site | - |
| Hallett\textsuperscript{18} | AAA | - | 2.5\% per patient | - |
| | | | 1.25\% per anastomosis | - |
| Mii\textsuperscript{24} | Global | 2.5\% per patient | - | - |
| Kalman\textsuperscript{25} | AAA | 12.8\% per patient | 7.5\% per patient | 5.3\% per patient |
| Biancari\textsuperscript{1} | AAA | - | 2.9\% per patient/aortic site | - |

PAAA, para-anastomotic aneurysm of the abdominal aorta; AAA, abdominal aortic aneurysm; AOD, aortoiliac obstructive disease; PAOD, peripheral artery obstructive disease.
et al.\(^4\) estimated overall incidences of PAAA of 0.8%, 6.2% and 35.8% after 5, 10 and 15 years, respectively.

PAAAs are challenging for surgeons because of the technical difficulties, which are caused by intense para-aortic fibrosis, encountered during surgical preparation of the aorta near the PAAA, which increase the risk of both iatrogenic damage to vessels and organs and peri-operative complications. Technical aspects and surgical variables are, therefore, crucial in determining a patient’s operative risk.\(^6\) Moreover, patients with PAAA are 8-10 years older than at the time of the primary graft and their co-morbidities (particularly cardiac and pulmonary) are expected to be worse.\(^6\) These factors contribute to making the operative risk for PAAA higher than that for primary aortic surgery, which is strictly related to co-morbidities.

It is clear that a ruptured or seriously complicated PAAA increases the operative risk, therefore, emergency treatment is associated with a higher surgical risk than elective management. In agreement with reports in the literature (Table 6), there was a very high mortality rate (66.6%) in the subgroup of our patients treated as emergencies, whereas the mortality rate was only 5% in the electively treated subgroup \(p = 0.005\), two-tailed Fisher’s exact test). On the other hand, we found no significant differences in early post-operative morbidity rates between elective and emergency cases. Furthermore, no significant differences were found in mortality rates between patients undergoing localized and extensive surgery \(p = 0.09\), 2-tailed Fisher’s exact test). In conclusion, elective surgery of an uninfected PAAA carries a higher operative risk than that of primary grafting, however, it is acceptable in terms of mortality, when compared with the risk associated with emergency treatment.\(^3,6,33\)

It is clear from the foregoing that early diagnosis is extremely important. An uninfected PAAA may evolve silently over a long period. Symptoms are absent or unusual. When present, they are non-specific and compatible with numerous differential diagnoses. Clinical signs are usually difficult to detect due to the retroperitoneal location of PAAA,\(^4\) therefore, imaging methods are necessary to make a diagnosis. The easiest, least invasive and most repeatable examination is color Doppler ultrasound, which reveals an abdominal aortic dilatation,\(^34\) although its nature and extension cannot always be defined. Angio-CT is always diagnostic and is the method of choice in order to define the extension and size of the aneurysm, the involvement of visceral arteries, relationships with adjacent structures and determination of the exact location of the PAAA.

### Table 6. Mortality and Morbidity in Literature

| PAAA                    | Mortality                        | Morbidity                        |
|-------------------------|----------------------------------|----------------------------------|
| Treiman\(^4\)           | False 8% in election 67% in emergency | 36% in election                  |
| Curl\(^7\)              | False & True 24% global 17% in election 67% in emergency 9% in election with infrarenal reconstructions 28% in election with suprarenal reconstructions | -                                |
| Allen\(^9\)             | False & True 21% global 17% asymptomatics | 73% global 33% asymptomatics    |
| Hagino\(^5\)            | False & True 0%                  | 27%                              |
| Coselli\(^2\)           | True / Recurrent 12.2% in-hospital 16.7% with rupture | 17% global intra-operative 53% global post-operative 14% intra-operative in election 50% post-operative in election 32% intra-operative in emergency 70% post-operative in emergency 58% with localized intervention 44% with extensive intervention |
| Mulder\(^6\)            | False 7.6% global 4.5% in election 24% in emergency 5.3% with localized intervention 8.5% with extensive intervention | 17% global intra-operative 53% global post-operative 14% intra-operative in election 50% post-operative in election 32% intra-operative in emergency 70% post-operative in emergency 58% with localized intervention 44% with extensive intervention |
| Matsumura\(^3\)         | False and True 5.1% in election 88% in emergency | -                                |
| Locati\(^3\)            | False and True 37.5% global in-hospital 14% asymptomatics 70% symptomatics | -                                |

PAAA, para-anastomotic aneurysm of the abdominal aorta.
organs and all the anatomical and morphological data necessary to plan the treatment strategy. In emergency cases, angio-CT is the method of choice to make the diagnosis and collect as much anatomical data as possible and also as quickly as possible. It is essential that esophagogastroduodenoscopy is performed prior to angio-CT in patients with acute or chronic gastrointestinal bleeding and a history of infrarenal aortic grafting in order to evaluate the origin of hematemesis or melena.

All patients treated with abdominal aortic grafting must be followed-up with standardized, long-term evaluations, including clinical examination, color Doppler ultrasound and angio-CT. After a learning curve, we now perform clinical and abdominal ultrasound examinations 2 months after the intervention and then annually. Angio-CT should initially be reserved for patients whose ultrasound findings indicate the need for this investigation and then performed in all cases 5 years after the intervention, when the incidence of graft infection and PAAA begins to become significant. Such a periodic, life-long surveillance program should enable PAAA to be treated in elective conditions and with an acceptable interventional risk.

Uninfected PAAA in symptomatic patients must always be treated. The indications to treat asymptomatic true PAAA are strictly related to the dimensions and the rate of growth of the aneurysm, as for generic abdominal aortic aneurysms: a sac diameter of 5 cm or more or twice the graft or normal aortic diameter. Many authors emphasize the importance of aggressive management of uninfected false PAAA: the reported incidence of rupture ranges from 8 to 55%, particularly when a surveillance program is not routinely performed. With regards to asymptomatic false PAAA, the diagnosis itself represents an indication to treat because of the fast, unpredictable evolution and higher incidence of rupture compared to that of true PAAA (in our population, all emergency cases were false PAAA). A pseudoaneurysm consists of a fibrotic capsule whose resistance to mechanical stress is lower than that of the native aortic wall, therefore, rupture is more frequent than rupture of a true PAAA and relatively independent of diameter. A watch-and-wait strategy seems to be justified only in electively manageable patients with a short life-expectancy and/or high surgical/anesthetic risk. However, careful evaluation of the risk/benefit ratio should be made when dealing with small false PAAA.

Endovascular treatment of uninfected PAAA is based on the larger experience reported with infrarenal aortic aneurysms and has advantages that contribute significantly to reducing peri-operative morbidity and mortality. Our database includes information on only 3 patients who underwent endovascular treatment, but it is clear that this less invasive approach reduces complications and shortens time spent in the hospital by eliminating the above described technical/surgical risks. There were no peri-operative deaths in our subgroup of patients who were managed with an endovascular procedure and the complications described were minor events with a weak relationship with the procedure. The endovascular option for uninfected PAAA, however, has some limitations: the absence of an infrarenal neck long enough for an endoanastomosis (if considering a pararenal PAAA); the high risk of a type II endoleak from the patent caudal aorta or iliac arteries after proximal end-to-side reconstruction; the management of residual hypogastric vasculature; tortuosity and poor flexibility of iliac arteries/grafts due to periprosthetic fibrosis; persistent pressure of the PAAA on adjacent organs after effective exclusion of the aneurysm; graft infection; the absence of long-term results completely validating AAA endovascular treatment; and the failure of endograft/Dacron fixation. In particular, failed fixation of tube stent-grafts inside old Dacron, causing endotension and sac rupture has been documented, therefore, bifurcated endografts appear to be a more durable solution for exclusion of a PAAA. On the other hand, it has been suggested that tube aortic cuffs can sequentially be used in an overlapping configuration to create customized endografts and overcome the difficulties described above. The current use of fenestrated/branched endografts (which we have not yet experimented) has extended the indications for endovascular treatment to thoraco-abdominal and juxtarenal/pararenal aortic aneurysms. Such configurations which do not interfere with aortic side branches have been found useful in endovascular treatment of uninfected PAAA, resulting in significant reduction of major adverse events and death in the majority of cases lacking an infrarenal neck. Further trials are, however, necessary to validate the long-term efficacy of these newly available devices, which are associated with considerable incidences of early endoleak and in-stent stenosis in fenestration sites.

In accordance with published data, our results confirmed the need for life-long clinical and imaging surveillance after aortic grafting; follow-up methods must be the cheapest and least invasive that enable early diagnosis of uninfected PAAA. This complication develops silently, and is difficult to repair and associated with a high mortality rate if treated as an emergency because of life-threatening complications. Elective surgical repair is advocated in order to minimize mortality and morbidity rates. Endovascular therapy with new generation devices is less invasive, bypasses surgical technical difficulties, and may represent an important option to combine safe treatment with a reduction of peri-operative adverse events.

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REFERENCES

1. Biancari F, Ylönen K, Anttila V, Juvonen J, Romsø P, Tatjana J, et al. Durability of open repair of infrarenal abdominal aortic aneurysm: a 15-year follow-up study. J Vasc Surg 2002;35:87-93.

2. Crawford ES, Saleh SA, Babb JW 3rd, Glaser DH, Vaccaro PS, Silverson A. Infrarenal abdominal aortic aneurysm: factors influencing survival after operation performed over a 25-year period. Ann Surg 1981;193:699-709.

3. Matsumura JS, Pearce WH, Cabellon A, McCarthy WJ 3rd, Yao JS. Reoperative aortic surgery. Cardiovasc Surg 1999;7:614-21.

4. Tréliman GS, Weaver FA, Cosson DM, Foran RF, Cohen JL, Levin PM, et al. Anastomotic false aneurysms of the abdominal aorta and the iliac arteries. J Vasc Surg 1988;8:268-73.

5. Hallett JW Jr, Marshall DM, Pettersson TM, Gray DT, Bower TC, Cherry KJ Jr, et al. Long-term survival and late complications after repair of ruptured abdominal aortic aneurysms. J Vasc Surg 1999;28:813-9; discussion 819-20.

6. Edwards JM, Teefey SA, Zierler RE, Kohler TR. Intraabdominal para-anastomotic aneurysms after infrarenal aortic aneurysm surgery. J Vasc Surg 1998;25:442-31; discussion 443-6.

7. Curl GR, Faggioni GL, Stella A, D’Addato M, Ricotta JJ. Aneurysmal change at or above the proximal anastomosis after infrarenal aortic grafting. J Vasc Surg 1992;16:855-9; discussion 859-90.

8. Gautier C, Borie H, Lagneau P. Aortic false aneurysms after prosthetic reconstruction of the abdominal aorta. J Vasc Surg 1993;7:8-13.

9. Signorelli M, Gaeta A, De Nale A, Tosini S, Ranucci M, Tealdi DG. Pseudoaneurismi anastomotici: una complicazione nella chirurgia vascolare ricostruttiva. Atti del XXVI World Congress of the International College of Surgeons, Milan 03-09 July 1988. In: Free papers printed in full. Bologna: Monduzzi Editore, 1988.p.561-5.

10. Gaylis H. Anastomotic false aneurysms of the abdominal aorta and the iliac arteries. J Vasc Surg 1988;8:268-73.

11. Plate G, Hollier LA, O’Brien P, Pairolero PC, Cherry KJ. Recurrent aneurysms and late vascular complications following repair of abdominal aortic aneurysms. Arch Surg 1985;120:590-4.

12. Illig KA, Green RM, Ouriel K, Rigs P, Bartos S, DeWeese JA. Fate of the proximal aortic cuff: implications for endovascular aneurysm repair. J Vasc Surg 1997;26:492-90; discussion 499-501.

13. Lisko DA, Ernst CB. Natural history of the residual infrarenal aorta after infrarenal abdominal aortic aneurysm surgery. J Vasc Surg 1998;27:805-11; discussion 811-2.

14. Bhalla SP, Sivakumar RS, Ranganathan MS, Prabhu M, Arivazhagan A, et al. Aortic pseudoaneurysms: a continuing late complication of vascular reconstructive procedures. Arch Surg 1986;121:314-7.

15. Kalman PG, Rappaport DC, Merchant N, Clarke K, Johnston KW. The value of late computed tomographic scanning in identification of vascular abnormalities after abdominal aortic aneurysm repair. J Vasc Surg 1999;29:442-50.

16. Taylor LM Jr, Van Kolken RJ, Baur GM, Porter JM. Precise diagnosis of aortic anastomotic aneurysm by computed tomographic scan. Arch Surg 1981;116:1209-11.

17. Guinet C, Buyijn JN, Ghossain MA, Mark AS, Jardim M, Fourmestraux J, et al. Aortic anastomotic pseudoaneurysms: US, CT, MR, and angiography. J Comput Assit Tomogr 1992;16:182-8.

18. Busuttil SJ, Goldstone J. Diagnosis and management of aortoenteric fistulas. Semin Vasc Surg 2001;14:302-11.

19. Bianchi P, Dalainas I, Ramponi F, dell’Aglio D, Casana R, Nano G, et al. Late gastrointestinal bleeding after infrarenal aortic grafting: a 16-
40. Kalman PG. What are the long-term results of conventional open surgical repair of abdominal aortic aneurysms? Acta Chir Belg 2003;103:197-202.

41. Melliere D, Berrahal D, Becquemin JP, Desgranges P, Cavillon A. [False anastomotic aneurysms after aorto-femoral prosthesis. Detection, prevention and treatment.] J Mal Vasc 1996;21:158-64.

42. Yuan JG, Marin ML, Veith FJ, Ohki T, Sanchez LA, Suggs WD, et al. Endovascular grafts for noninfected aortoiliac anastomotic aneurysms. J Vasc Surg 1997;26:210-21.

43. Liewald F, Kapfer X, Görich J, Halter G, Tomczak R, Scharrer-Pamler R. Endograft treatment of anastomotic aneurysms following conventional open surgery for infrarenal aortic aneurysms. Eur J Vasc Endovasc Surg 2001;21:46-50.

44. Magnan PE, Albertini JN, Bartoli JM, Ede B, Valerio N, Moulin G, et al. Endovascular treatment of anastomotic false aneurysms of the abdominal aorta. Ann Vasc Surg 2003;17:365-74.

45. Faries PL, Won J, Morrissey NJ, Briggs VL, Cadot H, Carroccio A, et al. Endovascular treatment of failed prior abdominal aortic aneurysm repair. Ann Vasc Surg 2003;17:43-8.

46. van Herwaarden JA, Waasdorp EJ, Bendermacher BL, van den Berg JC, Teijink JA, Moll FL. Endovascular repair of paraanastomotic aneurysms after previous open aortic prosthetic reconstruction. Ann Vasc Surg 2004;18:280-6.

47. Zhou W, Bush RL, Bhama JK, Lin PH, Safaya R, Lumsden AB. Repair of anastomotic abdominal aortic pseudoaneurysm utilizing sequential AneuRx aortic cuffs in an overlapping configuration. Ann Vasc Surg 2006;20:17-22.

48. Anderson JL, Adam DJ, Berce M, Hartley DE. Repair of thoracoabdominal aortic aneurysms with fenestrated and branched endovascular stent grafts. J Vasc Surg 2005;42:600-7.

49. O’Neill S, Greenberg RK, Haddad F, Resch T, Sereika J, Katz E. A prospective analysis of fenestrated endovascular grafting: intermediate-term outcomes. Eur J Vasc Endovasc Surg 2006;32:115-23.

50. Greenberg RK, West K, Pfaff K, Foster J, Skender D, Haulon S, et al. Beyond the aortic bifurcation: branched endovascular grafts for thoracoabdominal and aortoiliac aneurysms. J Vasc Surg 2006;43:879-86; discussion 886-7.

51. Adam DJ, Berce M, Hartley DE, Anderson JL. Repair of juxtarenal para-anastomotic aortic aneurysms after previous open repair with fenestrated and branched endovascular stent grafts. J Vasc Surg 2005;42:997-1001.

52. Verhoeven EL, Muhs BE, Zeebregts CJ, Tieljui IF, Prins TR, Bos WT, et al. Fenestrated and branched stent-grafting after previous surgery provides a good alternative to open redo surgery. Eur J Vasc Endovasc Surg 2007;33:84-90.