Repair of Giant Midline Abdominal Wall Hernias: “Components Separation Technique” versus Prosthetic Repair

Interim Analysis of a Randomized Controlled Trial

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Abstract

Background: Reconstruction of giant midline abdominal wall hernias is difficult, and no data are available to decide which technique should be used. It was the aim of this study to compare the “components separation technique” (CST) versus prosthetic repair with e-PTFE patch (PR).

Method: Patients with giant midline abdominal wall hernias were randomized for CST or PR. Patients underwent operation following standard procedures. Postoperative morbidity was scored on a standard form, and patients were followed for 36 months after operation for recurrent hernia.

Results: Between November 1999 and June 2001, 39 patients were randomized for the study, 19 for CST and 18 for PR. Two patients were excluded perioperatively because of gross contamination of the operative field. No differences were found between the groups at baseline with respect to demographic details, co-morbidity, and size of the defect. There was no in-hospital mortality. Wound complications were found in 10 of 19 patients after CST and 13 of 18 patients after PR. Seroma was found more frequently after PR. In 7 of 18 patients after PR, the prosthesis had to be removed as a consequence of early or late infection. Reherniation occurred in 10 patients after CST and in 4 patients after PR.

Conclusions: Repair of abdominal wall hernias with the component separation technique compares favorably with prosthetic repair. Although the reherniation rate after CST is relatively high, the consequences of wound healing disturbances in the presence of e-PTFE patch are far-reaching, often resulting in loss of the prosthesis.
wall reconstruction because of the technical difficulties, the high morbidity, and the relatively high recurrence rate associated with these procedures. However, many patients with large hernias have invalidating complaints such as bulging of the abdominal wall, chronic wounds, immobility, and back pain, necessitating surgical treatment.

The lack of sufficient tissue requires the insertion of prosthetic material or transposition of autologous material to bridge the fascial gap. Reconstruction using pre-peritoneally placed prosthetic material is still the most frequently applied method of reconstruction.1 The increased risk of infection in case of wound complications is a relative contraindication against the use of prosthetic materials. Moreover, interposition of either peritoneum or greater omentum between the bowel and the prosthesis is often impossible, which is another reason to avoid the use of prosthetic material.

In 1990 Ramirez, Ruas, and Dellon introduced the "components separation technique (CST)" to bridge the fascial gap without the use of prosthetic material.2 The technique is based on enlargement of the abdominal wall surface by separation and advancement of the muscular layers. In this way, defects of up to 20 cm at the waistline can be bridged. Retrospective series report promising results, but no prospective study has been published until now.3–12 It was the aim of this prospective study to compare the results of prosthetic repair with CST in patients with giant abdominal wall hernias that cannot be closed primarily. The primary endpoint of the study was reherniation; secondary endpoints were operation time and postoperative wound complications. In the present report the results of an interim analysis are presented.

PATIENTS AND METHODS

Adult patients (18–80 years) with an incisional hernia after midline laparotomy with a cranio-caudal length of at least 20 cm that could not be closed primarily, in whom the repair could be performed under clean or clean-contaminated conditions, and who were not using corticosteroid therapy were asked to participate in the study. Patients with perioperative gross contamination of the operative field were excluded from the study. After written informed consent the patients were randomized between CST and prosthetic repair by the data center of the Radboud University Nijmegen Medical Centre using envelope, the day before operation.

Fully trained abdominal wall surgeons who had done at least five procedures of both techniques before the start of the study performed the operations. H.v.G. and R.P.B. performed supervision in centers not having this expertise. Before each procedure, a preoperative chest x-ray was made. Demographic data, co-morbidity (COPD, cardiovascular disease, or diabetes), body mass index, condition of the skin, size of the hernia at the time of the operative procedure, kind of anesthesia, operation time, perioperative blood loss, postoperative ICU stay, analgesia use, complications, hospital stay, and follow-up were recorded on a standard form. The study protocol was reviewed and approved by the institutional ethics commissions of all the participating hospitals. All patients gave written informed consent after receiving a thorough explanation of the study.

Operative Technique

Standard thrombosis (Nadroparine 2,850 IE) and antibiotic prophylaxis (cefazoline 3·1 g and metronidazole 3·500 mg) were started preoperatively. After induction of anesthesia (combined general and epidural, if possible) and disinfection of the skin with iodine tincture, an adhesive drape was applied on the skin, if possible. The abdomen was entered via a midline laparotomy or at the lateral edge of the graft if the bowels were covered with a split skin graft. Adhesions between the ventral abdominal wall and the intra-abdominal viscera were cut, after which the length and width of the defect were measured.

Components Separation Technique (CST Group)

The component separation technique was performed as described in detail in former publications.2,12,13 Briefly, the skin and subcutaneous fat are dissected free from the anterior rectus sheath and the aponeurosis of the external oblique muscle (Figure 1A). The aponeurosis of the external oblique muscle is transected longitudinally about 2 cm lateral from the rectus sheath, including the muscular part that inserts on the thoracic wall, which extends at least 5–7 cm cranially of the costal margin (Figure 1B). The external oblique muscle is separated from the internal oblique muscle as far laterally as possible (Figure 1B). The posterior rectal sheath is separated from the rectus abdominis muscle if tension-free closure is impossible (Figure 1C). The fascia is closed in the midline with a running polydioxanone suture (PDS-loop, Johnson & Johnson, Ltd.) of at least 4 times the length of the incision. The skin is closed over at least two closed suction drains.
Prosthetic Repair (e-PTFE Group)

The skin and subcutaneous tissue are mobilized from the underlying fascia of the rectus abdominis muscle. As a consequence all epigastric perforating arteries supplying the overlying skin are separated. After adhesiolysis, a 20 x 30 cm, 1.5-mm-thick e-PTFE patch (Gore-Tex dual mesh plus with holes, W. L. Gore and associates Inc., Flagstaff, AZ, USA) is shaped in size and implanted intra-abdominally as underlay with an overlap of at least 4 cm to the aponeurosis, as described elsewhere.\textsuperscript{14} The mesh is placed intra-abdominally as an underlay and is sutured under slight tension to the ventral abdominal wall using a double row of interrupted sutures of e-PTFE 1/0 (Gore-Tex 1/0, W. L. Gore and associates Inc., Flagstaff, AZ, USA) that passed the rectus abdominis muscle. The prosthesis is implanted with the microporous side facing the intra-abdominal viscera and the macroporous side facing the fascia. (As a consequence of the large size of the hernias, the fascia could not be closed over the prosthesis in any of the patients in our series.) After implantation, the skin is closed over at least two closed suction drains.

Postoperative Care

Antibiotic prophylaxis, cefazoline 3 x 1 g and metronidazole 3 x 500 mg was started preoperatively and continued for the first 24 h postoperatively. All patients have epidural anesthesia if possible. Wounds were inspected on a daily basis with respect to hematoma, seroma, skin necrosis, and wound infection. Hematoma was defined as an accumulation of blood in the operative field for which a surgical intervention (puncture or drainage) was needed; seroma as an accumulation of fluid in the operative field for which an intervention (puncture or drainage) was needed in case of mechanical or physical limitations. Skin edge necrosis was defined as necrotic loss of full thickness skin for which surgical intervention was needed.

The wound was scored on a daily basis according to CDC criteria, as follows:\textsuperscript{15} grade 1: normal wound, grade 2: erythema and swelling, grade 3: purulent effluent; or grade 4: open wound. Drains were removed after 5 days or if production was less than 50 ml/24 h.

The thorax was examined daily by physical examination, and a routine x-ray of the thorax was performed on the second and seventh days after the operation, to detect pneumonia and atelectasis. No specific instructions were given to the patients after operation and patients had no restriction of physical activity except heavy lifting. Follow-up was done in the outpatient clinic at 3, 6, 12, 24, and 36 months after operation. At each visit a physical examination was done to diagnose recurrent hernia. Ultrasonography or computed tomography (CT) scanning was performed on indication, especially to detect small recurrences.

Figure 1. Operative technique of the “components separation technique.” 1 = rectus abdominis muscle; 2 = external oblique muscle; 3 = internal oblique muscle; 4 = transversus abdominis muscle; 5 = posterior rectal sheath. A. Dissection of skin and subcutaneous fat. B. Transaction of aponeurosis of external oblique muscle and separation of internal oblique muscle. C. Mobilization of posterior rectal sheath and closure in the midline. Adapted from Bleichrodt et al.\textsuperscript{13}, with permission of Elsevier.

Statistical Analysis

Patients were analyzed as per intention to treat. Hernia recurrence-free survival was compared using the Kaplan-Meier methods according to the intention-to-treat principle.

Power Analysis

Type I and II errors were set to 0.05 and 0.1, respectively. The minimum relevant difference in reherniation between groups was set to 30%, in advantage of CST.
Accordingly, a minimum of 84 patients was required (two
groups of 42 patients). An interim analysis was planned
to evaluate the results of the trial after inclusion of 40
patients.

Differences between groups were analyzed using the
Fisher exact test for demographic data, preoperative and
perioperative data, wound complications, reoperation,
and reherniation (Table 1).

RESULTS

Between November 1999 and June 2001, 39 patients
were included in the study and were operated on by one
of 5 surgeons. Two patients were excluded from the study
because of gross contamination during operation. Nineteen
patients, 6 women and 13 men, were randomized to
the CST group: reconstruction using the components
separation technique. The mean age of these 19 patients
was 53.9 years (range: 33–73 years). Eighteen patients,
6 women and 12 men, were randomized to the e-PTFE
group (prosthetic repair). Their mean age was 58.7 years
(range: 42–82 years).

In the CST group, closure of the fascia was accom-
plished in 18 of the 19 patients (Figure 2). In one patient
the abdominal wall hernia was too large and had to be
repaired using a combination of the CST and prosthetic
repair. In the e-PTFE group the procedure was suc-
cessful in 17 of the 18 patients. In one patient the
abdominal wall hernia was too large and was recon-
structed using a combination of prosthetic repair and
CST. No differences were found between the groups with
respect to demographic data (Table 1), co-morbidity,
length and width of the defect, skin coverage, anesthesia,
blood loss, and ICU stay (Table 1). All operations were
performed without major intraoperative complications,
except for the two excluded patients with gross periop-
erative contamination. The operation time for prosthetic
repair was significantly longer as compared with the
components separation technique ($p < 0.001$, Fisher
exact test) (Table 1). This is mainly due to the time-
consuming fixation of the patch to the fascia with a double
row of single sutures.

Postoperative Mortality and Morbidity

There was no 30-day mortality. Major wound compli-
cations were found in 10 of the 19 patients in the CST
group: wound infection ($n = 3$), skin necrosis ($n = 2$),
hematoma ($n = 1$). Four patients developed seroma;
these were not associated with the aforementioned
complications.

Major wound complications were found in 13 of the 18
patients in the e-PTFE group: wound infection ($n = 2$),
skin necrosis ($n = 3$), hematoma ($n = 1$). Both wound
infection and skin necrosis ultimately resulted in loss of
the prosthesis (Table 1). Seven patients developed a
seroma. In two of these patients seroma puncture was
performed to prevent spontaneous evacuation via the
midline wound; this resulted in infection and, ultimately,
loss of the patch. Seven patches were removed after a
median period of 94 days (range: 30–262 days). In the
cases where the prosthesis was removed, the abdominal
wall defect was reconstructed using CST.

Pulmonary complications were found in 4 patients in
the CST group and 2 in the e-PTFE group (not significant,
Fisher exact test) (Table 1).

Reherniation

Follow-up was complete in all patients. Four patients in
the CST group died before the end of the follow-up period
5, 9, 10, and 12 months after the operation from unrelated
causes. Two had a reherniation at the time of death. Of
the remaining 15 patients, 8 had a reherniation. Recur-
cences occurred after a mean period of 7 months (range:
0.5–12 months). Recurrences were all located in the
midline in the upper abdomen and were small. Two pa-
tients underwent reconstruction of their recurrence. One
patient in whom the reconstruction was performed with a
combination of CST and prosthetic bridging had a recurrent hernia at the edge of the prosthesis (Figure 3).

Prosthetic Repair

One patient in the e-PTFE group died 6 months after the operation from an unrelated cause. Of the remaining 17 patients, 7 had an infected prosthesis that had to be removed. The abdominal wall defect was then reconstructed using CST repair. Four other patients had a small recurrent hernia after prosthetic repair, without complaints. Recurrences occurred after a mean period of 22 months (range: 6–36 months). None of these four patients underwent reoperation for their recurrence (Figure 3).

DISCUSSION

The present study is the first randomized controlled trial comparing different techniques to repair giant midline

| Table 1. Study characteristic of patients with prosthetic repair or components separation technique |
|-------------------------------------------------|-------------------|-------------------|-------------------|
| Group 1: prosthetic repair | Group 2: components separation technique | Significance |
| Age (mean) | 58.7 (range: 42–82) | 53.9 (range: 33–37) | (t-test) | NS |
| Gender (women/men) | 6/12 | 6/13 | (Fisher exact test) | NS |
| BMI | 28.7 (range: 21.5–39.6) | 28.2 (range: 23.9–38.7) | (t-test) | NS |
| Defect (median) (cm) | 25 (range: 20–30) | 25 (range: 20–33) | NS |
| Width | 17 (range: 9–30) | 15 (range: 7–25) | NS |
| Skin (n) | | | (t-test) | |
| Intact, full thickness | 14 | 12 | NS |
| Intact, split skin | 4 | 7 | NS |
| Anesthesia (n) | | | (Fisher exact test) | |
| General | 3 | 5 | NS |
| Epidural and general | 15 | 14 | NS |
| Operative time (min) | 183 (range 135–254) | 113 (range 63–175) | p < 0.001 | (t-test) |
| Blood loss (ml) | 420 (range 100–900) | 289 (range 50–1000) | NS |
| ICU stay | | | (Mann-Whitney U-test) | |
| Patients (n) | 6 | 3 | NS |
| Time (days) | 2 (range: 1–6) | 5 (range: 1–10) | NS |
| Pulmonary complications (n) | 2 | 4 | NS |
| Pneumonia | 2 | 1 | NS |
| Atelectasis | 0 | 3 | NS |
| Analgesia | | | (t-test) | |
| Epidural (days) | 2.4 (range: 0–5) | 2.4 (range: 0–6) | NS |
| Morphine (days) | 3.3 (range: 0–8) | 3.6 (range: 0–10) | NS |
| Wound complication (n) | 1 | 1 | NS |
| Hematoma | 7 | 4 | NS |
| Seroma | 3 | 2 | NS |
| Skin necrosis | 2 | 3 | NS |
| Infected mesh (n) | 7 | 0 | (t-test) | |
| Reoperation (in OR) for wound complications (n) | 7 | 2 | p = 0.05 |
| Recurrence (n) | 11 | 10 | NS |

BMI: body mass index; ICU: intensive care unit; NS: not statistically significant; OR: Operation room.
hernias and the first prospective trial regarding the "component separation technique." Although our series is relatively small, the results suggest that repair of giant abdominal wall defects with the component separation technique compares favourably with prosthetic repair, because wound infection in patients in whom a prosthetic repair was performed had major consequences, resulting in removal of the prosthesis in 7, whereas wound infection in patients after CST had only minor consequences.

Disturbed wound healing frequently complicates repair of large abdominal wall hernias. Wound complications such as hematoma, seroma, skin necrosis, and infection are reported in 12%–67% of patients after CST and in 12%–27% after prosthetic repair. Wound complications are associated with the extensive dissection needed in both procedures, which are often performed after intra-abdominal catastrophes. The risk is further increased by the long duration of the operative procedure and the need to mobilize the skin in dividing the epigastric perforating arteries (Figure 4). This endangers the blood supply of the skin, because then it solely depends on the intercostal arteries, which may have been damaged during former operations by introduction of drains, or by stoma construction and other procedures needed in patients with intra-abdominal sepsis. Wound complications in our series were rather frequent. Although they are mentioned in most other publications about CST, the method of follow-up is mentioned in only one other study from our own group.

Loss of the prosthesis may also be associated with the choice of the prosthetic material used. Several materials have been developed for hernia repair. In the present series only patients with giant and often complex hernias were included. In the majority of these patients the peritoneum or greater omentum was not available to interpose between the prosthesis and the intra-abdominal viscera. Therefore, an e-PTFE dual patch was used to bridge the fascial gap. The expanded-PTFE dual patch has significantly better mechanical properties than polypropylene-mesh. It is a soft pliable microporous material that causes no mechanical trauma to the viscera. The micropores on both sides of the patch are too small to allow ingrowth of fibrocollagenous tissue, thus preventing fibrous adhesions on the visceral side of the patch. Lack of ingrowth results in insufficient anchorage of the patch to the adjacent fascia, however, and this is a major disadvantage of e-PTFE patches. The patch should be placed as underlay with an overlap of at least 4 cm and fixed to the aponeurosis with a double row of single sutures.

The e-PTFE patch is prone to infection because of its hydrophobic characteristics. To reduce the infection risk, the e-PTFE patch used is impregnated with silver salts and chlorhexidine, which both have anti-microbial properties and work synergistically. Moreover, antibiotic prophylaxis was given to all patients and an adhesive drape was applied to the skin. Nevertheless, 40% of our patients had an early or late infection resulting in removal of the patch. In a recent experimental study in rats with a large abdominal wall defect, it was found that impregnation with silver salts resulted in an aggravated inflammatory response around the patch and an increased reherniation rate. This observation may explain the increased risk for seroma formation, which is associated with prosthetic loss in this study. Some patients (n = 3, 16%) were operated under clean-contaminated condition,
which means they had an accidental bowel lesion during adhesiolysis without gross contamination. We suspect that most surgeons still place a prosthetic patch for abdominal wall reconstruction in these situations, which is supported by some small series in the literature.

In our opinion polypropylene, which is still the most widely used material for hernia repair, is contraindicated because of its propensity for inducing extensive visceral adhesions and occasional fistula formation. If large areas of polypropylene mesh are exposed, scar contraction will result in wrinkling of the polypropylene mesh, causing mechanical irritation, which promotes infection and carries the risk of mesh erosion into the skin or the intestine. If the polypropylene mesh cannot be covered with full-thickness skin, chronic infection and sinus formation will ultimately result in loss of the mesh. Therefore the results probably would not have been better if polypropylene mesh or polypropylene mesh based prosthesis was used.

Recurrent hernia still is a major problem. The only randomized controlled trial comparing open suture and mesh repair of small ventral hernias was reported by Luijendijk et al., reporting recurrence rates of 46% and 23%, respectively, after a follow-up of 36 months and 63% and 32%, respectively, after a follow-up of 75 months. In retrospective studies recurrence rates of 25%–63% in suture repair and 8%–25% in mesh repair are reported. Tension-free repair of incisional hernia is a prerequisite to prevent recurrence. In CST a tension-free repair was accomplished. In the literature recurrence rates of 0%–28% have been reported for CST, although how follow-up was accomplished is not well documented in most series. But it seems impossible to have a reherniation rate, in series of large abdominal wall defects, that is far below the reherniation rate of reconstruction of small abdominal wall defects in a well performed randomized controlled trial.

Despite the high recurrence rate in the present study and our retrospective study, CST remains an attractive technique for repair of giant ventral hernias. Most recurrent hernias are small and asymptomatic and need no further treatment. In addition, the functional and cosmetic results are good and patients were satisfied.

In a recent other study in 39 patients undergoing CST repair for heavily contaminated abdominal wall defects, similar results were found with respect to complications and reherniation rate (36%). All but one patient indicated satisfaction with the result when compared to their situation before operation. In that study, postoperative quality of life was assessed using the SF 36 questionnaire. When compared to the general population, patients had an average score or higher on pain, vitality, social functioning, and role limitations (emotional problems); the score was below average on physical functioning, role limitations (physical problems), in general health perception, and in mental health.

On the basis of the interim analysis, the trial was discontinued because the frequency of wound complications resulting in subsequent prosthetic loss was unacceptably high. Because underlay repair necessitates transection of the perforating epigastric arteries in patient with prosthetic repair it was expected that this complication could not be prevented, whereas CST remains possible if the epigastric perforators are spared. Impregnation of the e-PTFE patch with silver salts and chlorhexidine might have contributed to this. Recently, a prospective randomized controlled trial has started comparing CST with CST + preperitoneal polypropylene mesh support, combining the advantages of CST and prosthetic repair.

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