A Randomized Controlled Trial Comparing Packing with Nonpacking of Perianal Abscess Cavity

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Abstract

Introduction: Anorectal abscess is a potentially debilitating condition and one of the common anorectal conditions encountered in practice. Anorectal abscesses are defined by the anatomic space in which they develop and are more common in the perianal and ischiorectal spaces. Traditional management of perianal abscess involves early incision, drainage, curettage and packing of the residual cavity. Cavity packing and dressings are problematic in that they cause significant discomfort and require multiple visits to change the packing until healing. An alternative to the traditional approach is to perform adequate incision and drainage and then to allow healing without the use of cavity dressings.

Objective: To compare the effects of packing with nonpacking of the perianal abscess cavity on patient discomfort and wound healing and subsidiary evaluation of the clinical outcomes.

Materials and Methods: This single centre randomized controlled trial was carried out on 50 patients who were operated upon for perianal abscess in Combined Military Hospital, Dhaka during the period of July 2014 to June 2016. Patients were randomly assigned to receive either packing or nonpacking treatment through allocation by sealed envelope. The packing (control) group was instructed to report to a single nursing staff for subsequent dressing with packing of the residual cavity. The non-packing group was discharged with a superficial protective dressing; they did not undergo wound dressing but managed their own wounds until follow-up. Outcome measures were time of healing, abscess recurrence, fistula formation and post operative pain.

Results: A total of 54 patients were enrolled (4 lost in follow-up): 24 in the packing and 26 in the nonpacking arm. The two groups were comparable in terms of age and gender distribution, type, size of the abscess, duration of symptoms and length of follow-up. Healing in the non-packing group was faster compared to that of the packing group: mean 24.08 days versus 34.13 days (P=0.000). The rate of abscess recurrence was similar (P=0.664). Post operative fistula rates were similar (P=0.623). Pain scores appeared less in the nonpacking arm and statistically significant (7.25 vs. 4.24, P=0.000).

Conclusion: Small size of the study population was the limitation of this study. In order to obtain a higher level of evidence, an adequately powered multicentre based prospective randomized controlled trial is required to definitely address the question of packing of the cavity and its beneficial outcome following incision and drainage in the management of perianal abscess.

Key-words: Perianal abscess, Non-packing, Packing, Anorectal disease.
Traditional management of perianal abscess involves early incision, drainage, curettage and packing of the residual cavity. The rationale for cavity dressings is initially to provide surgical haemostasis and then to prevent skin closure over the cavity, allowing healing by secondary intention. Cavity packing and dressings, however, are problematic in that they cause significant discomfort and require multiple visits to change the packing until healing has occurred. There is currently no established evidence for this practice however; regular post operative packing can be painful.

An alternative to the traditional approach is to perform adequate incision and drainage and then to allow healing without the use of cavity dressings. The current guidance from The American Society of Colon and Rectal Surgeons suggests that with an adequately sized elliptical incision, post operative wound packing is usually not necessary. In the UK, however, this has not been incorporated into standard practice, and in the absence of sufficient evidence that it is both safe and effective, most UK Surgeons continue to pack the cavity until healing is achieved, especially for perianal abscesses. There is currently no robust evidence to guide clinical practice. The aim, therefore, was to examine the effects of this approach on patient discomfort and wound healing with subsidiary evaluation of the clinical outcomes. A randomized, prospective trial was conducted to compare the conventional treatment of perianal abscess with that of incision and drainage without cavity dressings.

**Materials and Methods**

This was a single-centre randomized controlled trial. This study was carried out on 50 patients who were operated upon for perianal abscess in Combined Military Hospital (CMH), Dhaka. The recruitment period was from July 2014 to June 2016.

All patients, adults aged 20 years or above, presenting with anorectal abscess was included in this study. Exclusion criteria included patients under the age of 20 years or unwilling to give consent and associated other conditions like known fistulae, diabetes mellitus, crohn’s disease, immune suppression, malignancy or other underlying causes. Patients with recurrent abscesses where the initial abscess drainage was considered inadequate (if the skin was not opened sufficiently to allow drainage of the abscess cavity) and those who underwent primary fistulotomy during drainage were also excluded from the study.

The patient recruitment was performed on first post operative day prior to any dressing change. Patients were randomly assigned to receive either packing or nonpacking through allocation by sealed envelope. An informed written consent was obtained from each patient willing to participate in the study.

Patients were treated with incision and drainage of the abscess cavity under general or spinal anaesthesia. The abscess was deroofed (involving an elliptical excision of perianal skin over the abscess cavity of sufficient length to drain the entire cavity and any extensions) to allow free drainage of the residual cavity. All patients were given a haemostatic pack intraoperatively, so the surgeon was blinded to the allocation of study group and had their dressing changed on first post operative day by the nursing staff.

On discharge, the packing (control) group were instructed to report to a single nursing staff of the hospital for subsequent dressing with packing of the residual cavity. Patients in the non-packing group were discharged with a superficial protective dressing to absorb any discharge from the cavity and protect the open wound. These patients were advised to have sitz bath, keep the area as clean and dry as possible. They did not undergo wound dressing but managed their own wounds until follow-up.

Data were collected on patient demographics, characteristics of the abscess (size and type), length of hospital stay, duration of symptoms before presentation, time for cavity healing, recurrent abscess or fistula formation, pain scores and length of follow-up. For the purposes of this study, abscesses were classified as superficial (superficial/submucosal) or deep (ischiorectal/ intersphincteric). Pain scoring was achieved via a standard Visual Analog Scale (VAS) for pain, at the first outpatient visit, in which subjects were asked to score their average pain over the previous 2 weeks, rather than daily discomfort levels. The primary end-point was time for cavity healing. Secondary end-points were abscess recurrence, fistula formation and pain score at two week using a standard visual analogue scale (VAS).

The predefined follow-up period for each patient was until the abscess cavity had healed completely. The patients were then followed up for a minimum 6-month period for abscess recurrence or fistula formation. All patients were reviewed at two weekly intervals in the outpatient clinic until healing occurred. Healing was defined as the cavity being closed and the skin completely re-epithelialized.
For patients who failed to attend the outpatient clinic, telephone interviews were conducted to determine time of healing (no longer requiring dressings, no further discharge), pain scores, and other data.

**Results**

Sixty patients presenting with perianal abscess were assessed for eligibility for the trial. Fifty-five patients were eligible for inclusion and 54 were enrolled. Patients were divided into the packing (control) group and the non-packing (intervention) group. Twenty-seven patients were allocated to each group. As block randomization was used, the end result involved equal patients in each group. Three subjects from packing group and one from non-packing group were lost in follow-up. All follow-up reviews were conducted in person. The flow of participants through each stage of the trial is described in Fig-1.

All continuous data were tested for normal distribution by nonparametric statistical method and analyzed accordingly using Mann-Whitney U test. Fistula formation and abscess recurrence rate were analyzed using Fisher’s exact probability test. Both tests were two-tailed with a significance level of 0.05. Analysis was performed using SPSS statistical software version 19 (SPSS Inc. Chicago, IL, USA).

The two groups were comparable on a demographic basis with regards to age and gender distribution. Characteristics of the abscess in terms of type (superficial or deep), size, duration of symptoms before presentation and length of follow-up were also similar between the groups. Demographic data and abscess characteristics are outlined in Table-I.
Table-I: Patient Demographics and Abscess Characteristics

<table>
<thead>
<tr>
<th></th>
<th>Packing (n=24)</th>
<th>Non-packing (n=26)</th>
<th>P value</th>
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<tbody>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median (Range)</td>
<td>37.5(21-65)</td>
<td>36.5(20-65)</td>
<td>0.331</td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>19</td>
<td>20</td>
<td>0.850</td>
</tr>
<tr>
<td>Female</td>
<td>05</td>
<td>06</td>
<td></td>
</tr>
<tr>
<td>Abscess Type No (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Superficial</td>
<td>16(66.7%)</td>
<td>18(69.2%)</td>
<td>0.848</td>
</tr>
<tr>
<td>Deep</td>
<td>8(33.3%)</td>
<td>8(30.8%)</td>
<td></td>
</tr>
<tr>
<td>Median Abscess Size</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(Range in mm)</td>
<td>25(25-35)</td>
<td>25(20-35)</td>
<td>0.437</td>
</tr>
<tr>
<td>Median Duration of Symptoms</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(Range in Days)</td>
<td>02</td>
<td>03</td>
<td>0.061</td>
</tr>
<tr>
<td>Median Length of Follow-up</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(Range in Week)</td>
<td>4(3-24)</td>
<td>3.5(14-90)</td>
<td>0.773</td>
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The mean time of healing (defined as complete epithelialization of the wound) was 34.13 (range, 16–110) days in the packing and 24.08 (range, 14–90) days in the non-packing group with significant P value (P=0.000). Post operative fistulas were detected at follow-up in eleven patients: six patients in the packing group (25%), and five (19.2%) in the non-packing group; this was not a significant difference (P=0.623). There was also no difference in abscess recurrence rates between the groups (3/24 in packing group vs 2/26 in non-packing group; P=0.664). The non-packing group reported less pain at 2 weeks post operatively. At two weeks, the median pain score in the non-packing group was 4.25 compared with 7.25 in the packing group (P=0.000). There were no differences in median length of stay (P=0.709). No adverse or unexpected events were seen in either group. Results are summarized in Table-II.

Table-II: Post operative Outcome

<table>
<thead>
<tr>
<th></th>
<th>Packing (n=24)</th>
<th>Non-packing (n=26)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Time of Healing Mean Days (Range)</td>
<td>34.13(16-110)</td>
<td>24.08(14-90)</td>
<td>0.000</td>
</tr>
<tr>
<td>Fistula Formation (%)</td>
<td>6(25%)</td>
<td>5(19.2%)</td>
<td>0.623</td>
</tr>
<tr>
<td>Recurrent Abscess (%)</td>
<td>2(8.3%)</td>
<td>2(7.7%)</td>
<td>0.664</td>
</tr>
<tr>
<td>Post Operative Pain Score at Two week Median (Range)</td>
<td>7.25(4-9)</td>
<td>4.25(1-9)</td>
<td>0.000</td>
</tr>
<tr>
<td>Length of Stay-Median Days(Range)</td>
<td>1(1-2)</td>
<td>1(1-2)</td>
<td>0.609</td>
</tr>
</tbody>
</table>

Discussion

Anorectal abscess is believed to originate from an infection in the anal glands. In 1880, Hermann and Desfosses demonstrated branching of the anal glands within the internal sphincter, submucosa and opening into the anal crypts. They were the first to suggest that infection in the anal glands results in extension of sepsis through the inter sphincteric space to the perianal tissues. Tucker and Hellwing demonstrated definitively that anal sepsis originates in the anal ducts which allows the infection to extend from the anal lumen into the wall of the anal canal.

The infection may extend between the internal and external sphincter, reach the anal verge to become a perianal abscess or it may rupture through the external sphincter and become an ischiorectal abscess. If the abscess extends cephalad in the rectal wall, a high intermuscular abscess will result and extension of abscess above the levators will produce a supralever abscess. A deep postanal abscess may extend to either or both ischiorectal fossae resulting in a horseshoe abscess.

The conventional management of perianal abscess often results in prolonged cavity dressings associated with increased cost, pain and inconvenience to patients with no evidence to support the practice. Alternative approaches to the surgical management of superficial abscesses have been investigated. Curettage and primary closure of the anorectal abscess cavity (30 cases) under antibiotic cover was first described by Ellis in 1951 who reported a high rate of primary healing (6.5 days). Ellis in 1960, reported a primary intention healing rate of 77 percent after primary suture of the curetted anorectal abscess cavity (151cases). Abraham et al reported a good result in terms of faster healing time and less time off work compared to packing of the abscess cavity. The primary healing rate was 78% but this approach was applied to soft tissue abscesses, perianal abscesses were not studied exclusively. Curetting of abscess cavities under antibiotic cover with primary closure by suturing, although advocated by Ellis and others, this approach has failed to gain wide popular acceptance.
De Pezzer catheter drainage is another alternative to traditional wound packing. In 1990, Kyle and Isbister retrospectively compared the use of a de Pezzer catheter with traditional incision and drainage. The authors indicated that the de Pezzer catheter was well tolerated and compared favorably with the traditional technique.

O’Malley et al undertook a randomized trial on the treatment of cutaneous abscesses (on the trunk, buttock and limbs) without cavity packing. Their results revealed that non packing did not cause any increased morbidity and the patients reported decreased pain scores. Koehler et al observed that cutaneous abscesses may be treated with incision and drainage alone thus avoiding pain and hospitalization. Kessler demonstrated in a randomized trial that wound packing does not impact failure or recurrence rates after simple incision and drainage, though there was no significant difference in pain scores. These information indicate that abscess cavities can be managed without the use of cavity dressings without an increase in complication rates. However, in many hospitals continued packing is recommended.

Tonkin et al conducted a pilot study in 2004 comparing two groups of patients presenting with perianal abscesses treated by incision and drainage of the abscess cavity with and without packing. Their results demonstrated that mean healing times (p=0.214), rate of abscess recurrence (p=0.61), post operative fistula rates were similar (P=0.38) and pain scores appeared much reduced in the non-packing arm, but did not attain statistical significance. They concluded that perianal abscesses can be managed safely without continued packing of the cavity. This study demonstrates reduced healing time and reduced post operative pain at 2 weeks. These findings were not demonstrated by Tonkin et al. This study also found that packing did not confer any protection with respect to risk of subsequent abscess recurrence and fistula formation. Perera et al demonstrated that healing in the non-packing group was faster when compared to that of the packing group (mean 26.8 days, P=0.047), the non-packing group reported less pain at 2 weeks post operatively (P=0.030) and there were no differences in recurrence rates between the groups (P=0.58). These findings are conforming to the present study.

A 2016 Cochrane database of systematic review demonstrated that it is unclear whether using internal dressings (packing) for the healing of perianal abscess cavities influences time to healing, wound pain, development of fistulae, abscess recurrence or other outcomes. Despite this absence of evidence, the practice of packing of abscess cavities is common. Given the lack of high quality evidence, decisions to pack may be based on local practices or patient preferences. Further clinical research is needed to assess the effects and patients’ experience of packing.

Conclusion
This study demonstrates that perianal abscesses can be treated safely and effectively with incision and drainage of the abscess cavity alone with no requirement for continued cavity packing while packing of the abscess offered no protection with regards to abscess recurrence and fistula formation. The non-packing of anorectal abscess cavity is likely to provide significant savings in terms of nursing resources and to reduce the patient discomfort associated with frequent cavity dressing changes. The authors acknowledge that it’s small population limits the study. In order to obtain a higher level of evidence, an adequately powered multicenter based prospective randomized controlled trial is required to definitely address the question if packing of the cavity is necessary or beneficial following incision and drainage in the management of perianal abscess.

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References


