Efficacy of Single Pre Operative Dose of Glucocorticoid (125 mg of Solumedrol Intravenous) in terms of Seroma Formation in Patients Undergoing Mastectomy with Axillary Clearance for Breast Cancer: A Randomized Controlled Trial

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Abstract

Objective: The study aimed to determine the efficacy of single dose of glucocorticoid (125 mg of Solumedrol intravenous) in terms of seroma formation after mastectomy in patients with carcinoma of breast.

Study design: Randomized controlled trial.

Place and duration of study: Study was conducted in the Department of General Surgery, Liaquat National Hospital Karachi, Pakistan from July 1 to Dec 31, 2010.

Patients and methods: Patients were randomly divided in two groups (study and control) each consisting of 30 patients. Randomization was done by opening of a sealed envelope which had a slip bearing the name of study medication (solumedrol or saline as placebo) to be administered. The study group received a single dose of inj 125 mg solumedrol IV half an hour prior to surgery by resident scrub in surgery. A similar procedure was applied to the control group and patients in controlled group were administered an equal volume of saline intravenously. After drain removal patients in both groups were observed for duration of 2 weeks for seroma formation. Detection of seroma formation was based on clinical grounds by absence of any fluid collection at mastectomy bed as detected by manual palpation. SPSS 10 was used for analysis.

Results: Seroma formation was observed in 66.7% (40/60) women 2 weeks post drain removal. Rate of seroma formation was significantly low in study groups than control groups (33.3% vs. 100%; p = 0.0001).

Conclusion: Single dose of steroid is efficacious in reducing the post mastectomy seroma formation.

Keywords
Post mastectomy seroma formation, Steroid and seroma, Factors for seroma formation, How to reduce seroma formation post mastectomy, Mastectomy, Seroma

Introduction

Seroma is a post operative fluid collection requiring one or more aspirations or subsequent drain placement following surgery. It is a common occurrence following surgery. The incidence rate of seroma formation following breast surgery is up to 30% to 92% [1]. It is often an ongoing problem after removal of the suction drain, and repeated skin puncture is necessary to remove the seroma. Seroma formation is most likely the result of the inflammatory response due to wound healing. Comparison of drain fluid to plasma ratios with known lymph to plasma values for biochemical parameters showed that this fluid is compositionally different from lymph, but is similar to inflammatory exudates [2]. In the seroma fluid several factors have been detected that support this assumption. These factors are: High levels of IgG, leucocytes, granulocytes, proteinases, pro-
teinases inhibitors, different kinds of cytokines (tPA, uPA, uPAR, PAI-1, PAI-2, IL-6, IL-1). The modulation of acute phase response may have important implications for patients with cancer undergoing surgery [3]. On the basis of this understanding, an inhibition of the inflammatory response might result in a decrease in seroma formation, and perhaps improve quality of life after mastectomy. Steroids inhibit the inflammatory response for example by inhibition of the cytokine function. It has been shown that a high single dose of steroid infusion (30 mg/kg solu-medrol) inhibits the normal IL 6 response after colon resection. While there are no studies on the use of solu-medrol in breast cancer, the use of triamcinolone in breast reconstruction surgery demonstrated than 55% did not have seroma formation whereas 95% developed seroma in those administered saline [4].

The exact etiology of seroma formation remains controversial. Several interventions have been reported with the aim of reducing seroma formation including the use of ultrasound scissors in performing quilting [5], buttress suture [6], lymphadenectomy [7], fibrin glue [8], fibrin sealant [9], bovine thrombin application [10], and altering surgical technique to close dead space [11].

The seroma formation after breast cancer surgery is independent of duration of drainage, compression dressing and other known prognostic factors in breast cancer patients except the type of surgery, i.e there is a 2.5 times higher risk of seroma formation in patients undergoing MRM compared to BP [12].

It is the possibility of decreased seroma formation with the use of steroids that we propose to undertake this study. It is proposed to use a single dose of 125 mg solu-medrol intravenously in this study.

Operational Definitions

Seroma: Presences of clear fluid filled pockets in the body post operatively; Efficacy: Efficacy will be deemed posture as when no seroma formation at the end of 4 weeks; Carcinoma Breast: Biopsy proven breast cancer lobular or infiltrative ductal.

Material and Methods

Study design: Randomized controlled trial; Setting: Study was conducted in the Department of General Surgery, Liaquat National Hospital Karachi, Pakistan from July 1 to Dec 31, 2010; Duration of study: 6 months, patients in each group were observed for seroma formation for 2 weeks after drain removal; Sample technique: Non probability purposive; Sample Size: Assuming the efficacy of solu-medrol 55% and saline 15% [12], level of significance 5% and power of test 90%, simple size required was 46 but we took 60 patient (considering lost to follow up) 30 patient in each group [4].

Sample Selection: Inclusion criteria:

• Women with primary breast cancer up to stage III, planned for a mastectomy with axillary dissection.
• Age over 25 years.
• Signed informed consent.

Exclusion criteria:

• Men
• Treatment with glucocorticoids within the last month before surgery, including inhalation products
• Pregnant.
• Severe heart disease
• Treatment with carbamazepine, phenytoin, Phenytoin, phenobarbital, rifampicin, salicylates and cyclosporine
• Uremia
• Diabetes
• Post neo-adjuvant and adjuvant chemotherapy.

Data collection procedure

Patients found eligible as per inclusion criteria were offered to participate in the study. Those giving informed consent were included. A proforma was filled for each patient to record. Mastectomy with axillary clearance was performed by consultant (with 15 years of experience) in breast surgery. Each patient enrolled in the study was eligible to be enrolled into either arm of the study following the opening of a sealed envelope which had a slip bearing the name of study medication (sOLUMEDROL or saline as placebo) to be administered. Most senior resident scrubbed in case was choosing the envelope and afterwards it was handed over to anesthetist who used to administer the randomized drug pre operatively to the patient during induction. The study group received a single dose of inj 125 mg solu-medrol IV half an hour prior to surgery by resident scrub in surgery. A similar procedure was applied to the control group and patients in controlled group were administered an equal volume of saline intravenously. Patients in both groups were followed postoperatively until 2 weeks after drain removal. An absence of clear fluid filled pocket in the mastectomy bed was documented in the proforma by a resident attending the OPD.

Data analysis procedure

Data was entered and analyzed by using statistical package for social sciences (SPSS) version 11.0 software. Frequencies and percentages were computed for categorical variables like age groups, stages of breast cancer, seroma formation and efficacy. Mean, standard deviations, 95% confidence interval, median with IQR were calculated for quantitative variables.
Results

A total of 60 women with primary breast cancer planned for a mastectomy with axillary dissection were included in this study. Patients were equally divided into two groups. The study group received a single dose of inj 125 mg solu-medrol IV half an hour prior to surgery and similar procedure was applied to the control group who were administered an equal volume of saline. Overall age distribution of the patients is presented in Figure 1. The average age of the patients was 52.32 ± 9.86 years (95% CI: 49.77 to 54.86) as shown in Table 1. Significant difference was not observed between group in average age (52.62 ± 8.42 vs. 52 ± 11.25; p > 0.05) as presented in Figure 2.

Regarding stages of carcinoma, 4 (7%) women were in stage 1A, 14 (23%) were in stage 2A 22(37%) were in stage 3A and 20 (33%) women were observed with...
Table 2: Comparison of seroma formation between groups.

<table>
<thead>
<tr>
<th>Seroma formation</th>
<th>Solumedrol n = 30</th>
<th>Normal saline n = 30</th>
<th>Total n = 60</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>10 (33.3%)</td>
<td>30 (100%)</td>
<td>40 (66.7%)</td>
<td>0.0001</td>
</tr>
<tr>
<td>No</td>
<td>20 (66.7%)</td>
<td>0 (0%)</td>
<td>20 (33.3%)</td>
<td></td>
</tr>
</tbody>
</table>

Chi-Square = 30, df = 1.

Table 3: Comparison of efficacy between groups.

<table>
<thead>
<tr>
<th>Efficacy</th>
<th>Solumedrol n = 30</th>
<th>Normal saline n = 30</th>
<th>Total n = 60</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>20 (66.7%)</td>
<td>0 (0%)</td>
<td>20 (33.3%)</td>
<td>0.0001</td>
</tr>
<tr>
<td>No</td>
<td>10 (33.3%)</td>
<td>30 (100%)</td>
<td>40 (66.7%)</td>
<td></td>
</tr>
</tbody>
</table>

Fisher’s exact test applied.

Table 4: Comparison of efficacy between groups for below and equal to 50 years of age.

<table>
<thead>
<tr>
<th>Efficacy</th>
<th>Solumedrol n = 14</th>
<th>Normal saline n = 15</th>
<th>Total n = 29</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>9 (64.3%)</td>
<td>0 (0%)</td>
<td>9 (31%)</td>
<td>0.0001</td>
</tr>
<tr>
<td>No</td>
<td>5 (35.7%)</td>
<td>15 (100%)</td>
<td>20 (69%)</td>
<td></td>
</tr>
</tbody>
</table>

Fisher exact test applied.

Stage 2B carcinoma as presented in Figure 3.

Seroma formation was observed in 66.7% (40/60) women 2 weeks post drain removal. Rate of seroma formation was significantly low in study groups than control groups (33.3% vs. 100%; p = 0.0001) as shown in Table 2. Comparisons of efficacy in term of seroma formation in women with primary breast cancer after mastectomy with axillary dissection are presented in Table 3. Efficacy of the treatments was significantly high in study group (Solumedrol) than control (66.7% vs. 0%; p = 0.0001). Similarly effectiveness of solu-medrol was significantly high with respect to age as presented in Table 4, Table 5 and Table 6 and it was also significantly effective in stage 2A, stage 3A and stage 2B as presented in Table 7, Table 8 and Table 9.

Discussion

The documented effect of triamcinolone as a potent anti-inflammatory agent could perhaps be sug-
In this study a total of 60 women with primary breast cancer planned for a mastectomy with axillary dissection were included. Patients were equally divided into two groups. The study group received a single dose of inj 125 mg solu-medrol IV half an hour prior to surgery and similar procedure was applied to the control group who were administered an equal volume of saline. The average age of the patients was 52.32 ± 9.86 years (95% CI: 49.77 to 54.86). Significant difference was not observed between group in average age (52.62 ± 8.42 vs. 52 ± 11.25; p > 0.05).

Regarding stages of carcinoma, 4 (7%) women were in stage 1A, 14 (23%) were in stage 2A 22 (37%) were in stage 3A and 20 (33%) women were observed with stage 2B carcinoma.

Seroma formation was observed in 66.7% (40/60) women. Rate of seroma formation was significantly low in study groups than control groups (33.3% vs. 100%; p = 0.0001). Efficacy of the treatments was significantly high in study group (Solumedrol) than control (66.7% vs. 0%; p = 0.0001). Similarly effectiveness of solu-medrol was significantly high with respect to age and it was also significantly effective in stage 2A, stage 3A and stage 2B.

While there are no studies on the use of solu-medrol (but easily available in our setup) in breast cancer, the use of triamcinolone (which has same

**Table 5:** Comparison of efficacy between groups for above 50 years of age.

<table>
<thead>
<tr>
<th>Efficacy</th>
<th>Solumedrol n = 16</th>
<th>Normal saline n = 15</th>
<th>Total n = 31</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>11 (68.8%)</td>
<td>0 (0%)</td>
<td>9 (31%)</td>
<td>0.0001</td>
</tr>
<tr>
<td>No</td>
<td>5 (31.3%)</td>
<td>15 (100%)</td>
<td>20 (69%)</td>
<td></td>
</tr>
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</table>

Fisher exact test applied.

**Table 6:** Comparison of efficacy between groups for stage 1A of breast cancer.

<table>
<thead>
<tr>
<th>Efficacy</th>
<th>Solumedrol n = 2</th>
<th>Normal saline n = 2</th>
<th>Total n = 4</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>1 (50%)</td>
<td>0 (0%)</td>
<td>1 (31%)</td>
<td>0.99</td>
</tr>
<tr>
<td>No</td>
<td>1 (50%)</td>
<td>2 (100%)</td>
<td>3 (75%)</td>
<td></td>
</tr>
</tbody>
</table>

Fisher exact test applied.

**Table 7:** Comparison of efficacy between groups for stage 2A of breast cancer.

<table>
<thead>
<tr>
<th>Efficacy</th>
<th>Solumedrol n = 7</th>
<th>Normal saline n = 7</th>
<th>Total n = 14</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>6 (85.7%)</td>
<td>0 (0%)</td>
<td>6 (42.9%)</td>
<td>0.005</td>
</tr>
<tr>
<td>No</td>
<td>1 (14.3%)</td>
<td>7 (100%)</td>
<td>8 (75.1%)</td>
<td></td>
</tr>
</tbody>
</table>

Fisher exact test applied.

**Table 8:** Comparison of efficacy between groups for stage 2B of breast cancer.

<table>
<thead>
<tr>
<th>Efficacy</th>
<th>Solumedrol n = 10</th>
<th>Normal saline n = 10</th>
<th>Total n = 10</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>4 (40%)</td>
<td>0 (0%)</td>
<td>4 (20%)</td>
<td>0.025</td>
</tr>
<tr>
<td>No</td>
<td>6 (60%)</td>
<td>10 (100%)</td>
<td>16 (80%)</td>
<td></td>
</tr>
</tbody>
</table>

Fisher exact test applied.

**Table 9:** Comparison of efficacy between groups for stage 3A of breast cancer.

<table>
<thead>
<tr>
<th>Efficacy</th>
<th>Solumedrol n = 10</th>
<th>Normal saline n = 10</th>
<th>Total n = 20</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>9 (81.8%)</td>
<td>0 (0%)</td>
<td>9 (40.9%)</td>
<td>0.0005</td>
</tr>
<tr>
<td>No</td>
<td>2 (18.2%)</td>
<td>11 (100%)</td>
<td>13 (59.1%)</td>
<td></td>
</tr>
</tbody>
</table>

Fisher exact test applied.
mode of action and duration like solu-medrol) in breast reconstruction surgery demonstrated that 55% did not have seroma formation whereas 95% developed seroma in those administered saline [12]. It is the possibility of decreased seroma formation with the use of steroids that we propose to undertake this study.

The small sample size of present study is a limitation and hence the power of the study is low. A number of questions remain unanswered and more research is needed to answer these.

Conclusion

Single dose of steroid is efficacious in reducing the post mastectomy seroma formation which helps in timely administration of chemotherapy and reduce the risk of infection, necrosis and sepsis and improve quality of life. Steroids may have applications in managing other seromas of similar etiology, such as those after abdominoplasty and hernia mesh repair.

References

3. Does a single steroid injection reduce the formation of post-mastectomy.