Extended Prophylactic Antibiotics for Mastectomy with Immediate Breast Reconstruction: A Meta-analysis

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INTRODUCTION

Breast cancer is one of the most common malignancies diagnosed in women and comprises about 18% of all female cancers. Surgery is the primary modality for the treatment of breast cancer, depending on tumor stage. An increasing number of breast cancer patients are opting for mastectomy with reconstruction for treatment.

Postmastectomy reconstruction can surgically restore the shape of the breast and provide breast cancer patients with psychological benefits. There are 2 types of postmastectomy reconstructions: autologous tissue flap and tissue expander/implant. Although autologous tissue provides the most lasting and natural outcomes, implant-based breast reconstruction is the more popular procedure, accounting for about 80% of postmastectomy reconstructions. Breast reconstruction can be divided by timing into immediate breast reconstruction (IBR) and delayed breast reconstruction (DBR). IBR, compared to DBR, offers a native inframammary fold and a pliable skin envelope for a more natural appearance. IBR also reduces psychological impact on patients and thus may be favored by some patients. However, compared to DBR, IBR is associated with greater risk of surgical site infection (SSI). SSI is defined as infection of the superficial incision, organ, and/or space after surgery. Accordingly, there are 3 categories of SSI: superficial incisional SSI, deep incisional SSI, and organ/space SSI. The rate of SSI is strongly related to the use of extended antibiotic prophylaxis. We aim to determine the effect of extended antibiotic prophylaxis on the incidence of SSI after IBR.

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associated with the type of surgical wound. The Centers for Disease Control (CDC) published a guideline in 1985,12 which classified surgical wounds into clean, clean/contaminated, contaminated, and dirty, with the SSI rate of 1%–5%, 3%–11%, 10%–17%, and over 27%, respectively.13 The occurrence of SSI can impact the postoperative recovery process and result in extra cost and rate of hospital readmissions.

Breast surgeries are classically categorized as clean, and according to the CDC and the Surgical Care Improvement Project, for breast surgical procedures, antibiotics should be discontinued within 24 hours after surgery.12 However, among breast surgeries, IBR with a tissue expander/implant is associated with higher SSI, with the average SSI rate ranging from 5% to as high as 35%.10,15,16

In the setting of IBR, several studies supported the use of extended prophylactic antibiotics to prevent SSI,17–19 but others stated that extended antibiotic usage could lead to systemic side effects and the development of resistant organisms.20–22 Thus, there is still no consensus regarding the extended usage, regimens, and timing of prophylactic antibiotics for mastectomy with IBR. This meta-analysis aims to determine the efficacy and safety of extended prophylactic antibiotics on SSI after mastectomy with IBR.

PATIENTS AND METHODS

Search Strategy

Our study followed guidelines published by the Centre for Reviews and Dissemination (CRD) and the Cochrane Collaboration and the Preferred Reporting Items for Systematic Reviews and Meta-Analyses criteria.23,24 A protocol for this systematic review was registered using Prospero (CRD42019127536).

We included patients undergoing mastectomy with IBR, or mixed types (IBR and DBR) if the study had separate outcomes for IBR and DBR groups. Both randomized controlled trials (RCTs) and observational studies were included. Studies that compare pre-, peri-, postoperative extended prophylactic antibiotics with standard of care were included. Studies with no comparison group were excluded.

Two reviewers independently assessed the title and abstract of articles identified by the search described earlier. Two reviewers applied the inclusion and exclusion criteria, and disagreements were resolved by reading the full text. A third reviewer examined the article and made the final decision if still undecided. The Preferred Reporting Items for Systematic Reviews and Meta-Analyses flowchart diagram of literature retrieval is shown in Fig. 1.

Data Extraction

Two reviewers independently extracted the data. Data were collected from each study: the first author, publication year, study design, number of procedures, type of IBR, antibiotics (ie, regimen name, dose, duration of treatment), and outcomes.

Quality Assessment for Included Studies

Two reviewers independently assessed the study quality using the Newcastle–Ottawa Scale.25 The Newcastle–Ottawa Scale is used to evaluate non-RCTs, with 1 version for case–control studies and the other for cohort studies. Both versions of the scale consist of 8 multiple-choice questions that address subject selection and comparability (of cases and controls in case–control studies and of cohorts in cohort studies) and the assessment of the outcome (in case–control studies) or exposure (in cohort studies).

Dealing with Missing Data

For those articles which met inclusion criteria but had incomplete data (ie, missing type of standard of care of comparison group, or no details of antibiotic use of extended antibiotics group), we emailed the corresponding author of that article. If corresponding authors did not respond after 3 weeks, the article was labeled as having incomplete data, and this would be mentioned in the results section.

Outcome Measures and Data Synthesis

All statistical analyses were performed by Review Manager V.5.3 (The Cochrane Collaboration, Software Update, Oxford, UK). We calculated the risk ratio (RR) at 95% confidence interval (CI) for each study, weighted by the number of events in each study. Statistical significance was defined as 2-tailed alpha < 0.05. Forest plots were generated for graphical presentations for clinical outcomes, and we used F statistics to define the heterogeneity of each study. Mantel-Haenszel method was used to conduct meta-analysis, and because F was >50%, we used the random-effects model instead of the fixed-effects model. We did subgroup analyses where the data were applicable.

Assessment of Heterogeneity

We assessed the heterogeneity between study results using the F statistics. The result is a percentage of total variation among studies due to heterogeneity. F is commonly divided into 3 categories—low, moderate, and high, with upper limits of 25%, 50%, and 75%, respectively.26

Sensitivity Analysis

We assessed the influence of a single study on the overall effect by sequentially removing 1 study at a time to test the robustness of the pooled results to further verify whether any study had an excessive influence on the overall results.

RESULTS

Results of the Search

A total of 597 articles were identified, of which 297 articles were excluded as they were ineligible publication types. After initial screening of the remaining 300 original articles, we further excluded 289 articles because they were irrelevant to our study topic or had incomplete data. The details of the search strategy are shown in Fig. 1.

Characteristics of Included Studies

A total of 11 studies (15,966 mastectomy procedures) were included, 10,688 in the extended antibiotics arm and
5,278 in the comparison arm. One was an RCT, and 10 studies were retrospective studies with 1 matched cohort study. The studies were published between 2009 and 2018, with sample sizes ranging from 112 to 7,443 mastectomies. The main characteristics of all 11 studies are presented in Table 1.

**Type of Procedures**

All procedures were unilateral or bilateral mastectomy with IBR. Immediate breast implant, tissue expander, and autologous flap (including latissimus dorsi flap and subpectoral flap) placement were described in 7, 3, and 2 studies, respectively.

**Timing and Types of Antibiotics**

Eight studies evaluated the usage of extended postoperative systemic prophylactic antibiotics compared to standard of care, where antibiotics are discontinued within 24 hours after breast surgery. However, the duration and types of antibiotics used vary between studies. In 5 of 8 studies, all patients were given pre- or perioperative antibiotics as baseline treatment, whereas the other 3 studies had incomplete data on antibiotic usage in their comparison groups. The study design was also different. Of the 8 studies, 1 study was an RCT, 2 studies were claims databases with a large sample size, and 5 studies were single-site retrospective studies.

Three studies evaluated the usage of postoperative topical antibiotics compared to no postoperative topical antibiotics. Two articles focused on the usage of topical mupirocin ointment and irrigation with both the extended antibiotics group and comparison arm using the same pre-/peri- and postoperative antibiotics. The third article used a novel antibiotic bead compared to comparison, where both arms had the same preoperative and irrigation antibiotics.

**Synthesis of Results**

**Incidence of SSI**

We found an overall average SSI rate of 5.99% in mastectomy procedures. Heterogeneity within the interventions used in the study prevented pooling of all the studies for analysis. Analysis of 8 studies comparing extended systemic antibiotics with standard of care found no statistical significance (RR = 0.80, 95% CI: 0.60–1.07, P = 0.13) (Fig. 2); as 1 RCT and
2 claims database studies are in the group of 8 studies, we also conducted other sensitivity analysis to check whether these studies may have affected the results (see further “Sensitivity Analysis” section). For the topical antibiotics group, analysis of the 3 studies comparing topical antibiotics with no topical antibiotics demonstrated statistically significant effect of antibiotics on reducing the incidence of SSI (RR = 0.26, 95% CI: 0.12–0.60, P = 0.001) (Fig. 3) However, this statistical significance has to be interpreted with caution, as more research is needed to confirm the findings. The use of topical mupirocin ointment and the use of novel antibiotic beads may not be generalizable to many other centers’ experiences.

### Wound Complications

Wound complications included hematoma, wound dehiscence, seroma, hematoma, and mastectomy flap necrosis. Four17,19,29,30 studies measured the incidence of wound complications, with 9,789 mastectomies on extended prophylactic antibiotics and 4,220 mastectomies on standard of care. Analysis of the 4 studies comparing extended prophylactic antibiotics with standard of care found no statistically significant effect on reducing the incidence of wound complications (RR = 0.89, 95% CI: 0.78–1.03, P = 0.12). The result is presented in Supplemental Digital Content 1, which displays a forest plot of wound complications subgroup analysis, [http://links.lww.com/PRSGO/B293](http://links.lww.com/PRSGO/B293).

### Hospital Readmission

Only 1 study measured the incidence of hospital readmission, with 6,049 mastectomies on extended prophylactic antibiotics and 1,349 mastectomies on standard of care. Analysis of the study comparing extended prophylactic antibiotics with standard of care found no statistical significance in reducing the rate of hospital readmission (RR = 1.22, 95% CI: 0.85–1.74, P = 0.28). The result is presented in Supplemental Digital Content 2, which displays forest plot of readmission subgroup analysis, [http://links.lww.com/PRSGO/B294](http://links.lww.com/PRSGO/B294).

### Table 1. Characteristics of Included Studies (n = 11)

<table>
<thead>
<tr>
<th>Study</th>
<th>Study Design</th>
<th>Type of Reconstruction</th>
<th>No. Operations</th>
<th>Intervention Group Antibiotics</th>
<th>Intervention Group SSI (%)</th>
<th>Control Group Antibiotics</th>
<th>Control Group SSI (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Avashia et al17</td>
<td>Retrospective</td>
<td>IBR TE</td>
<td>138</td>
<td>Postoperative &gt; 24 h</td>
<td>8/119 (6.7%)</td>
<td>Standard of care</td>
<td>6/19 (31.6%)</td>
</tr>
<tr>
<td>Hunsicker et al19</td>
<td>Retrospective</td>
<td>IBR implant</td>
<td>555</td>
<td>Postoperative for 96 h</td>
<td>6/316 (1.9%)</td>
<td>Standard of care + irrigation once</td>
<td>14/219 (6.4%)</td>
</tr>
<tr>
<td>Clayton et al18</td>
<td>Retrospective</td>
<td>IBR implant</td>
<td>250</td>
<td>Postoperative &gt; 24 h</td>
<td>21/116 (18.1%)</td>
<td>Standard of care</td>
<td>46/134 (34.3%)</td>
</tr>
<tr>
<td>Goh et al20</td>
<td>Retrospective</td>
<td>IBR TE, latissimus dorsi, subpectoral</td>
<td>240</td>
<td>Postoperative &gt; 24 h</td>
<td>12/145 (8.3%)</td>
<td>Standard of care</td>
<td>2/95 (2.1%)</td>
</tr>
<tr>
<td>Kenna et al27</td>
<td>Retrospective</td>
<td>IBR implant</td>
<td>127</td>
<td>Irrigation + antibiotic beads</td>
<td>1/8 (1.47%)</td>
<td>Standard of care</td>
<td>7/59 (11.86%)</td>
</tr>
<tr>
<td>McCullough et al22</td>
<td>Retrospective</td>
<td>IBR implant</td>
<td>378</td>
<td>Postoperative &gt; 24 h</td>
<td>24/290 (12%)</td>
<td>Standard of care</td>
<td>24/178 (13.5%)</td>
</tr>
<tr>
<td>Murray et al28</td>
<td>Retrospective</td>
<td>IBR TE</td>
<td>200</td>
<td>Topical ointment</td>
<td>0/23 (0%)</td>
<td>Standard of care</td>
<td>10/177 (5.65%)</td>
</tr>
<tr>
<td>Olsen et al29</td>
<td>Retrospective</td>
<td>IBR implant, flap, both</td>
<td>5,938</td>
<td>Postoperative &gt; 24 h</td>
<td>240/3,305 (7.26%)</td>
<td>Standard of care</td>
<td>213/2,633 (8.09%)</td>
</tr>
<tr>
<td>Phillips et al30</td>
<td>RCT</td>
<td>IBR implant</td>
<td>112</td>
<td>Postoperative &gt; 24 h</td>
<td>11/50 (22%)</td>
<td>Standard of care</td>
<td>12/62 (19.95%)</td>
</tr>
<tr>
<td>Ranganathan et al31</td>
<td>Retrospective</td>
<td>IBR implant</td>
<td>7,443</td>
<td>Postoperative &gt; 24 h</td>
<td>166/6,049 (5.26%)</td>
<td>Standard of care</td>
<td>41/1,394 (3.45%)</td>
</tr>
<tr>
<td>Townley et al32</td>
<td>Retrospective</td>
<td>IBR implant</td>
<td>605</td>
<td>Postoperative &gt; 24 h</td>
<td>9/297 (3.03%)</td>
<td>Standard of care</td>
<td>11/308 (3.6%)</td>
</tr>
</tbody>
</table>

TE, tissue expander.

![Fig. 2. Forest plot of studies that compared systemic antibiotics to standard of care (n = 8). SOC, standard of care. RR, risk ratio. M-H, Mantel-Haenszel.](http://links.lww.com/PRSGO/B293)
Quality Assessment for Included Studies

We created a quality assessment figure based on the Risk of Bias Tool found in the Cochrane Handbook for Systemic Reviews of Interventions, and we presented the percentages of risk in each of the 9 domains. High-quality responses were marked “low risk.” The result is shown in Supplemental Digital Content 3, which displays risk of bias,

Sensitivity Analysis and Publication Bias

Funnel plot analysis disclosed no asymmetry around the axis, which means that publication bias was not detected (Fig. 4). No significant results were identified in the “leave one out” sensitivity test.

Further Sensitivity Analysis to Understand Heterogeneity across Studies

As there were substantial differences in the types of studies within the 8 studies that investigated extended antibiotics on SSI, we conducted 2 further analyses. First, we noted that 2 studies using claims database had 88.5% of the weight (Fig. 2). We analyzed extended systemic antibiotics with and without the 2 claims database studies, resulting in 8 and 6 studies (Fig. 2). In Fig. 2, RR was 0.80 (95% CI: 0.60–1.07, I² = 61%). In Supplemental Digital Content 5, which displays a forest plot of extended systemic prophylactic antibiotics (without RCT study), RR was 0.77 (95% CI: 0.56–1.06, I² = 65%). Overall, the RCT did not change the results of the 8 studies. We note that given the nature of the study, theoretically the RCT should have been given more weight, but our predefined statistical method did not provide allowances for ad hoc increases in statistical weight.

DISCUSSION

For patients who are opting for mastectomy with reconstruction as treatment, IBR, especially implant-based IBR, has become a common procedure to restore the shape of the breast and improve psychological well-being. Breast surgery is historically thought of as a “clean” procedure, and for clean surgical procedures, the CDC calls for the discontinuation of perioperative prophylactic antibiotics within 24 hours. In recent years, some have advocated breast surgeries as “clean-contaminated” procedures, noting the breast microbiome and bacteria presence on normal breast implants, and of contamination of breast implants even when precautions are taken. However, this may be more complicated than applying a blanket label for all breast surgeries, because the SSI rates after different breast surgeries have varied widely in the published literature, ranging from 0.8% to 26%. Among all types of procedures, mastectomy with implant-based IBR has a 2-fold increase in SSI incidence compared with mastectomy.

Fig. 3. Forest plot of studies that compared topical antibiotics to standard of care (n = 3). SOC, standard of care. RR, risk ratio. M-H, Mantel-Haenszel.

Fig. 4. Funnel plot of included studies (n = 11). SE, standard error. RR, risk ratio.
I intake into 3 categories—low, moderate, and high, with upper estimation of variability among different studies. Thus, meta-analysis of heterogeneity is only 1 component of a wider evaluation of what constitutes an SSI.26 Thus, having clarity of what makes an SSI would be very important for any prospectively designed study. Disagreement on what constitutes an SSI is a commonly encountered problem when comparing articles discussing infection rates, as it may be field specific. We would like to engage stakeholders and experts in the area of breast reconstruction to potentially form a consensus for prospective studies that will improve evidence-based practices.

The third limitation is the type of studies included. Ten of 11 studies were retrospective studies. Since there was only 1 RCT conducted within the included studies, more well-designed RCTs should be conducted to demonstrate the effect of different regimens of prophylactic antibiotics on the SSI rate of IBR. Appropriate prophylactic antibiotic protocols should be tested. Finally, a better reporting system of the types of SSI, antibiotic regime/dosage/duration, and other complications should be used for future studies.

Future Directions

Given the paucity of prospective studies on this important topic, well-designed studies are sorely needed. However, there are many prior considerations that go into a well-designed prospective study for this particular question. Although we try not to prescribe particular rules here, the following questions (among others) are important to consider. Should patients with tissue necrosis be excluded? Should cases done by surgeons who leave completely different flap thicknesses be grouped together? Should reconstructions going under the muscle be compared to reconstructions done above the muscle? Should small areas of necrosis count the same as larger areas? Should the experience of the plastic surgeon be assessed? Should drain removal be based on drainage volume or based on duration, and if so, how much? Should patients who are discharged from hospital with a drain be grouped separately? It has been noted that these factors, while important in predicting SSI, are never available from retrospective chart reviews. Importantly, regarding antibiotic duration, the timepoints of antibiotics for 24 hours only, antibiotics until drain removal (which can vary significantly), and antibiotics for a certain duration after drain removal are essential to study and compare. Even if the sample size is relatively low, such well-controlled prospective studies will be valued. We rally surgeons in this field to consider starting prospective studies on this.

CONCLUSIONS

From our systematic review and meta-analysis, we conclude that, at this point, in part due to the lack of large prospective studies and in part due to the heterogeneity of interventions, there are insufficient data to suggest that extended antibiotics reduce the risk of SSI in patients undergoing mastectomy with IBR. Moreover, broad-spectrum antibiotics may significantly influence the normal gastrointestinal flora and lead to unfavorable clinical consequences, such as Clostridium difficile–related pseudo-membranous colitis and antibiotic resistance. Therefore, we appeal for RCTs that test if there is improved efficacy and safety of extended prophylactic antibiotics on IBR. In particular, focus should be put on the choice of antibiotic regimens, the treatment duration, and a standardized clinical criterion for SSI evaluation.
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REFERENCES

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