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(Article begins on next page)
Acellular Dermal Matrix in Reconstructive Breast Surgery: Survey of Current Practice among Plastic Surgeons

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Background: Acellular dermal matrices (ADMs) in plastic surgery have become increasingly popular particularly for breast reconstruction. Despite their advantages, questions exist regarding their association with a possible increased incidence of complications. We describe a collective experience of plastic surgeons’ use of ADMs in reconstructive breast surgery using an internet-based survey.

Methods: Members of the American Society of Plastic Surgeons were recruited through voluntary, anonymous participation in an online survey. The web-based survey garnered information about participant demographics and their experience with ADM use in breast reconstruction procedures. After responses were collected, all data were anonymously processed.

Results: Data were ascertained through 365 physician responses of which 99% (n = 361) completed the survey. The majority of participants were men (84.5%) between 51 and 60 years (37.4%); 84.2% used ADM in breast reconstruction, including radiated patients (79.7%). ADM use was not favored for nipple reconstruction (81.5%); 94.6% of participants used drains, and 87.8% administered antibiotics postoperatively. The most common complications were seroma (70.9%) and infection (16%), although 57.4% claimed anecdotally that overall complication rate was unchanged after incorporating ADM into their practice. High cost was a deterrent for ADM use (37.5%).

Conclusions: Plastic surgeons currently use ADM in breast reconstruction for both immediate and staged procedures. Of those responding, a majority of plastic surgeons will incorporate drains and use postoperative antibiotics for more than 48 hours. (Plast Reconstr Surg Glob Open 2015;3:e381; doi: 10.1097/GOX.0000000000000148; Published online 24 April 2015.)
Increased interest in ADM use for reconstructive breast surgery has mirrored the introduction of new products. Initially, ADM was reported for use in secondary breast deformities, such as contracture and rippling, but has since evolved to address other shortcomings of implant-based reconstruction. ADMs have been described as the most significant innovation impacting prosthetic breast reconstruction in recent years. This statement can be attributed to its numerous potential benefits, including improved aesthetic outcome, reduction in postoperative pain, and decreased operative time. Furthermore, it has been reported to provide better control of the mastectomy space, optimize implant positioning, allow for increased intraoperative expansion, and prevent superior migration of the implant. Prior studies have evaluated the outcomes of ADM use in breast reconstruction. Despite their advantages, there is literature implicating their association with an increased incidence of postoperative complications, particularly infection and seroma formation.

Although it is not common, ADMs can also be employed in delayed breast reconstruction provided there is an adequate degree of skin laxity. Moreover, ADMs have shown to be successful in patients with a variety of breast volumes. ADMs have proven less effective in patients with delayed reconstruction, in exposure to radiation, in those with a history of smoking, when vascularity to skin flaps has been compromised immediately following mastectomy, and in the morbidly obese. ADM in postmastectomy breast reconstruction in terms of the popularity of its use, the most commonly incorporated meshes, patient satisfaction, surgical outcomes, complications, and application in breast revision procedures. In addition, we sought to investigate the impact of ADM in the setting of drains, antibiotics, nipple reconstruction, and in previously radiated patients.

**METHODS**

**Recruitment**

A survey was created using http://www.surveymonkey.com. An invitation containing a generic link to the survey (that ensured anonymity and prevented tracking) was distributed by e-mail to 365 plastic surgeons who are members of the American Society of Plastic Surgeons. Participation was voluntary. After responses were collected, all data were anonymously processed. Only a small cohort of plastic surgeons who are members of the American Society of Plastic Surgeons was included owing to the difficulty in obtaining member e-mails (lack of an e-mail repository) and based on omission of e-mails that were either no longer in use by the plastic surgeon (referred to an administrative e-mail) or inactive, in which case the e-mail was not delivered (a delivery status notification failure was received). There were no incentives given for participation in this study, and those who did participate were completely random with no preselection based on practice patterns or ADM use.

**Survey**

The web-based survey gathered information about participant demographics, including gender, age, practice type, years in practice, and geographic setting. Participants were then asked if they utilized ADM in breast reconstruction. Plastic surgeons who did not use ADMs were asked about the use of biologic meshes in other procedures and questioned about their decision not to use ADM in breast procedures. For participants who used ADM in breast reconstruction, inquiries were made about the type of mesh, the reasons for their particular choice, application in either immediate or delayed setting, patient satisfaction with surgical outcomes, and the utilization of ADM in breast revision procedures. Furthermore, to better understand the scope of ADM application, inquiries were made regarding the use of ADM for nipple reconstruction, drain usage, administration of antibiotics postoperatively, and in previously radiated patients. Finally, participants provided insight into their experience with infection rates and complications following ADM use and their opinion of the literature concerning the current evidence pertaining to ADM use in plastic surgery.

**Statistical Analysis**

Statistical analyses were performed using SPSS software version 21.0 (SPSS Inc., Chicago, Ill.). Associations between the use of ADM (dependent variable) and different independent variables were determined using chi-square and Fisher’s exact tests. Two-sided P value of <0.05 was deemed statistically significant.
RESULTS

Patient Demographics

The data for this study were ascertained through 365 physician responses of which 99% \((n = 361)\) of the respondents completed the survey. Responses of plastic surgeons who did not complete the survey were excluded \((n = 4)\) (Fig. 1). The majority of participants were men \((84.5\%)\) who worked as solo practitioners in private practice \((51\%)\) in a large urban area \((60.1\%)\). Most plastic surgeons were between the ages of 51 and 60 \((37.4\%)\) and had been in practice for 11–20 years \((37.7\%)\). Participant demographics are summarized in Table 1.

Participants Who Did Not Use ADM in Breast Reconstruction

Of the 361 participants who completed the survey, 57 \((15.8\%)\) stated that they did not use ADM in breast reconstruction. When these 57 participants were queried as to whether they used ADM for any other procedures (such as abdominal wall reconstruction, head and neck reconstruction, burn surgery, lower limb coverage, and hand surgery), 42 \((73.7\%)\) claimed that they did not incorporate ADM into any aspect of their practice. The most reported reasons for this were due to cost \((n = 15)\), surgeon preference \((n = 10)\), and increased complications with previous experience \((n = 11)\). Only 15 of the 57 participants \((26.3\%)\) stated their use of ADM for other procedures, the most common of which were abdominal wall reconstruction \((n = 5)\) followed by extremity surgery, lower limb coverage \((n = 4)\), and

**Fig. 1.** Survey participant selection process.
hand surgery (n = 3). When asked about the decision not to incorporate ADM in breast procedures, most of the respondents attributed it to the absence of breast reconstruction in practice (n = 7), no clear indication of benefit (n = 3), and cost (n = 2).

Participants Who Use ADM in Breast Reconstruction

Three hundred four participants (84.2%) stated routinely using ADM in breast reconstruction. The majority of respondents in this group had been using ADM in breast procedures for the last 6–10 years (69.5%). The most popular mesh for use in practice was AlloDerm (LifeCell Corporation, Branchburg, N.J.; 71.6%) (Fig. 2). The main reason for its popularity was reported to be adequate long-term experience and AlloDerm being well described in the literature (68.1%). The data suggest that this group of participants chose to incorporate ADM in breast reconstruction to allow for better control of the implant pocket (81.4%), improved aesthetic outcomes (70.1%), a quicker expansion (43.9%), and being able to reduce the incidence of capsular contracture and breast deformities (40.2% and 25.6%, respectively).

The majority of responding plastic surgeons (81.5%) did not use ADM for nipple reconstruction. When asked about patient satisfaction, 86.4% of the surveyed plastic surgeon population reported that patients were satisfied with aesthetic outcomes; furthermore, 77.9% stated that patients rarely come back for further revisions. In the circumstance when a patient is unsatisfied and ADM has been used for reconstruction, the most commonly performed revision procedures include symmetry procedures (53.9%), fat grafting (19.5%), and capsulotomy/capsulectomy (17.9%).

Drains were used by 95% of respondents in conjunction with ADM in breast reconstruction; 81.5% reported using drains in all breast procedures involving ADM; typically, either 1 (45%) or 2 drains (49.6%) were used, and 57.5% of respondents stated to have left drains in for a longer period of time when they used ADM. Most participants (87.8%) routinely used antibiotics in the postoperative period; however, the number of days of antibiotic use varied from less than 5 days (31.5%), 6–10 days (45%), 11–14 days (15.8%), to more than 14 days (7.7%).

### Table 1. Demographic Characteristics of Respondents (n = 361)

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Total (%)</th>
<th>ADM Group (%)</th>
<th>Non-ADM Group (%)</th>
<th>P</th>
</tr>
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<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Male</td>
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<td>257 (84.5)</td>
<td>48 (84.2)</td>
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<td>47 (15.5)</td>
<td>9 (15.8)</td>
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<td>Age (y)</td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>&lt;30</td>
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<td>0 (0)</td>
<td>0 (0)</td>
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<tr>
<td>31–40</td>
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<td>20 (6.6)</td>
<td>0 (0)</td>
<td>0.05†</td>
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<tr>
<td>41–50</td>
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<td>104 (34.2)</td>
<td>14 (24.6)</td>
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<tr>
<td>51–60</td>
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<td>115 (37.8)</td>
<td>20 (35.1)</td>
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<td>&gt;60</td>
<td>88 (24.4)</td>
<td>65 (21.4)</td>
<td>23 (24.6)</td>
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<tr>
<td>Primary practice type</td>
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<td></td>
<td></td>
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<tr>
<td>Private solo</td>
<td>184 (51.0)</td>
<td>152 (50)</td>
<td>32 (56.1)</td>
<td>0.59*</td>
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<tr>
<td>Private group</td>
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<td>20 (6.6)</td>
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<td>1.00†</td>
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<td>Academic community</td>
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<td>7 (2.3)</td>
<td>1 (1.8)</td>
<td>1.00†</td>
</tr>
<tr>
<td>Academic university</td>
<td>49 (13.6)</td>
<td>40 (13.2)</td>
<td>9 (15.8)</td>
<td>0.59*</td>
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<tr>
<td>Practice setting</td>
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<tr>
<td>Large urban area</td>
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<td>174 (57.2)</td>
<td>43 (75.4)</td>
<td>&lt;0.01*</td>
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<tr>
<td>Small urban area</td>
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<td>12 (21.1)</td>
<td>&lt;0.01*</td>
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<tr>
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<td>16 (5.3)</td>
<td>2 (3.5)</td>
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<td>Years in practice</td>
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<td></td>
<td></td>
<td></td>
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<tr>
<td>&lt;5</td>
<td>5 (1.4)</td>
<td>5 (1.6)</td>
<td>0 (0)</td>
<td>0.60†</td>
</tr>
<tr>
<td>6–10</td>
<td>57 (15.8)</td>
<td>53 (17.4)</td>
<td>4 (7.0)</td>
<td>0.05†</td>
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<tr>
<td>11–20</td>
<td>136 (37.7)</td>
<td>118 (38.8)</td>
<td>18 (31.6)</td>
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</tr>
<tr>
<td>21–30</td>
<td>111 (30.7)</td>
<td>92 (30.3)</td>
<td>19 (33.3)</td>
<td>0.65*</td>
</tr>
<tr>
<td>≥30</td>
<td>52 (14.4)</td>
<td>36 (11.8)</td>
<td>16 (28.1)</td>
<td>&lt;0.01*</td>
</tr>
</tbody>
</table>

*Chi-square test.
†Fisher’s exact test.

n = 361; percentages shown have been calculated as a fraction of respective groups (ADM, n = 304; non-ADM, n = 57).
found that 79.7% of respondents had used ADM in previously radiated patients. Of these, 37.4% experienced no change in complication rates, an almost equal number of respondents suspected an increase or decrease in complication rate (24.3% versus 23%), and 15.3% were unsure. Those who reported an increase in complications implicated seroma (n = 45) and surgical site infection (n = 28), necessitating return to the operating room 11–30% of the time. The majority of participants (72.8%) used ADM in both immediate implant and staged reconstruction (tissue expander/implant). Regardless of the type of procedure, the most commonly reported complications were seroma (70.9%) followed by surgical site infection (16%) and wound dehiscence (9.4%) (Fig. 3). When asked whether ADM use in breast reconstruction contributed to an increased rate of infection, participants responded that it remained unchanged (57.4%); 26.4% of respondents reported an increase and 5.4% a decrease in the rate of infection. Finally, when questioned about their opinion of the literature with regard to current evidence for the use of ADM in plastic surgery, 38.7% of respondents suggested that the use of ADM is safe and effective in preventing complications while 28.4% reported an increase and 24.3% an equal number of respondents suspected an increase or decrease in complication rates (28.4% versus 24.3%). However, data about the demographics and its application in breast reconstructive surgery in the general plastic surgery population are less clear. In our study, the majority of responding plastic surgeons practicing in breast reconstruction (tissue expander/implant). Regardless of practice pattern, use ADM in breast reconstruction, which may suggest a practice pattern that newer graduates are learning and/or choose to use ADM in breast reconstruction at greater rates than surgeons who did not learn to use it during their training. This assertion is somewhat confirmed by the finding that 57.4% of respondents suspected that the rate of infection did not change with ADM. However, a careful review of recent evidence indicates that opinion is divided on whether or not the use of ADM is associated with a higher incidence of postoperative complications. A chart review of 41 patients (65 breasts) by Bindingnavele et al found extremely low complication rates with biologic mesh use in postmastectomy breast reconstruction: seroma in 3 patients, wound infection in 2 patients, and hematoma and expander removal in 1 patient each. This finding is corroborated in a study by Preminger et al who reported that AlloDerm did not increase the risk of postoperative complications. In contrast, Chun et al and Liu et al observed statistically significant increases in infection rate and seroma rate with AlloDerm use, respectively. Only 26.4% of our survey participants reported an increase in postoperative complications, the most common of which were seroma (70.9%), surgical site infection (16%), and wound dehiscence (9.4%). A systematic review by Ho et al showed higher likelihood of seroma and infection in prosthetic-based breast reconstructions using traditional musculofascial flaps, whereas Adetayo et al identified the most common complications as wound infection (16%), seroma formation (8%), and breast implant failure (6%).

![Fig. 3. Most common complications with ADM use in breast reconstruction.](image-url)
The ability of ADM to tolerate exposure to radiation is still being debated. In our study, 79.7% of participants claimed to have used ADM in previously radiated patients with most (37.4%) reporting no change in complication rate and equal numbers suspecting an increase (24.3%) or decrease (23%). Nahabedian reported that ADM is able to tolerate radiation exposure, demonstrating that the risk of infection did not vary with or without AlloDerm. Komorowska-Timek et al noted that AlloDerm reduced the rate of radiation-related inflammation. Colwell et al found that radiation therapy following stage 1 of tissue expander/implant-based reconstruction had a significantly lower complication rate than radiation therapy in the setting of breast conserving therapy. On the other hand, Spear et al and Salzberg et al observed 11-fold and 4-fold higher complication rates, respectively, in irradiated versus nonirradiated breasts. Despite these high complication rates, Ayeni et al in their review reported that compared with plain tissue expander reconstructions, ADM-assisted tissue expander reconstruction seemed to have better resistance to radiation or at least have similar complication rates.

No consensus exists on antibiotics use following breast reconstructive surgery. Despite indications that there is no benefit in patients who receive treatment for more than 24 hours, a majority of plastic surgeons (87.8%) routinely use antibiotics for 6–10 days (45%) in the postoperative period. Avashia et al demonstrated a significant decrease in the rate of infection when postoperative antibiotics were taken for at least 48 hours following implant-based breast reconstruction with ADM. In a series of 321 implant-based reconstructions of which AlloDerm was used in 75, Nguyen et al reported no variations in the readmission rates for intravenous antibiotics. However, development of infected fluid collections resulting in explantation was found to be significantly higher in the AlloDerm group compared with the control group. The use of drains was also prevalent in our surveyed population (94.6%) with just over half (57.5%) leaving drains in for a longer period of time when they used ADM. This finding is consistent with findings by Collis et al who reported drains to have remained in situ for a significantly longer duration when using ADM.

Few studies have implemented ADM to aid in reconstruction of the nipple-areola complex with the goal of improving nipple projection. The survey found that only 18.5% of respondents reported to have used ADM for nipple reconstruction. Although experience with ADM in nipple reconstruction is limited, results thus far may be promising. Garrazone and Lam demonstrated that AlloDerm use in a modified dermal flap pattern for 30 nipple reconstructions was a safe, reproducible, and easily performed approach for enhancing nipple projection. In contrast, a review of ADM use in nipple reconstruction by Israeli suggests limited success due to loss of nipple projection over time.
The most frequently used ADM is AlloDerm (71.6%), which is not surprising given it is the most commonly reported mesh for use in breast reconstruction in the literature. Moreover, 72.8% of surgeons used ADM in both immediate and staged breast reconstruction procedures owing to its many reported benefits. One major deterrent against ADM use in breast procedures is cost, which can range from $3536 to $4856 per breast; it was implicated as the main reason for not using ADM in practice at all by 37.5% of participants. Regardless, Salzberg found AlloDerm use in immediate reconstruction to be less costly than transverse rectus abdominis myocutaneous flap surgery and expander/implant reconstruction after mastectomy. Although ADM can be expensive, various reports have demonstrated that in the long-term, it is cost effective in breast reconstruction.

There are limitations to our study. Despite the number of plastic surgeons who completed the survey, only 57 did not use ADM for breast reconstruction. This finding was perhaps due to the voluntary sampling of participants which was utilized and may have resulted in bias toward ADM use. A small sample size may reduce the chances of detecting a true effect by overestimating it. Also, it may decrease the likelihood that a statistically significant result reflects a true effect. Therefore, this may be an incomplete assessment of the actual prevalence of plastic surgeons who do not use ADM in practice. Patient satisfaction with the aesthetic result was solely based on surgeon opinion, which may have also contributed toward bias with ADM use. Another limitation may be interpretation of the term “breast reconstruction” used in our survey, which some surgeons may have found to mean reconstruction following mastectomy only rather than also its use in aesthetic cases. Furthermore, the term “revision” may be have been interpreted differently by the study participants with some considering it to be further surgical intervention in the operating room and others deeming it a simple outpatient “touch-up.” Nevertheless, a majority of our respondents claimed that patients rarely came back for a revision maintaining consistency of our result. Finally, recall bias of the surveyed plastic surgeons may be a factor; however, this issue could potentially be overcome by our sample size and subgroup analysis excluding patients who did not use ADM for breast reconstruction.

CONCLUSIONS

Plastic surgeons use ADM in breast reconstruction for both immediate and staged procedures. Younger generations of plastic surgeons seem more willing to include ADMs in their practice. A majority of responding plastic surgeons incorporate drains and use postoperative antibiotics for more than 48 hours. ADM use for nipple reconstruction is not yet widely accepted. The occurrence of seroma, surgical site infection, and wound dehiscence are the most commonly implicated complications when ADM is incorporated necessitating return to the operating room. Despite this, a good number of respondents believe that overall infection rate remains unchanged. In addition, the majority of participants who reported use of ADM in previously radiated patients found that it did not contribute to a difference in complication rate. Most responding plastic surgeons believe that based on existing evidence, ADM use is safe and effective in preventing complications and that the data against its use are weak. The main deterrent against ADM use is its cost. In future studies, a larger participant population is needed to eliminate potential bias regarding ADM use.

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