



Characteristics of adhesion areas between the tissue expander and capsule in implant-based breast reconstruction

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Original Article

Background The use of anatomic implants has improved the aesthetic results of breast surgery; however, implant malrotation is an uncommon, but serious complication of these procedures. Nevertheless, little research has explored implant adhesion. In this study, we investigated adhesion between the expander and the capsule.

Methods Seventy-nine cases of immediate breast reconstruction via two-stage implant-based reconstruction performed between September 2016 and November 2017 were evaluated. Mentor CPX4 expanders were used in 14 breasts, and Natrelle expanders in 65. We analyzed areas of adhesion on the surfaces of the tissue expanders when they were exchanged with permanent implants. We investigated whether adhesions occurred on the cephalic, caudal, anterior, and/or posterior surfaces of the expanders.

Results Total adhesion occurred in 18 cases, non-adhesion in 15 cases, and partial adhesion in 46 cases. Of the non-adhesion cases, 80% (n = 12) were with Mentor CPX4 expanders, while 94.4% (n = 17) of the total adhesion cases were with Natrelle expanders. Of the partial adhesion cases, 90.7% involved the anterior-cephalic surface. The type of tissue expander showed a statistically significant relationship with the number of attachments in both univariate and multivariate logistic regression analyses ($P < 0.001$) and with total drainage only in the univariate analysis ($P = 0.015$).

Conclusions We sought to identify the location(s) of adhesion after tissue expander insertion. The texture of the implant was a significant predictor of the success of adhesion, and partial adhesion was common. The anterior-cephalic surface showed the highest adhesion rate. Nevertheless, partial adhesion suffices to prevent unwanted rotation of the expander.

Keywords Mammoplasty / Breast implants / Breast implantation / Rotation / Tissue expansion devices

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INTRODUCTION

The use of anatomical implants has led to improved aesthetic results following certain procedures, including breast recon-

struction and breast augmentation [1]. Accurately-shaped anatomic breast implants contain more filling material in the inferior pole of the implant than in the superior pole, in order to imitate the natural shape of the breast as closely as possible.

An uncommon, but seriously compromising complication that may occur following surgery is rotation of the implant. Because round implants are symmetrical, rotation does not typically cause them to appear malpositioned. However, rotation of anatomic implants can lead to the appearance of a misshapen breast [2,3].

In some studies, the authors have reported a rotation rate ranging from 0.9% to 14% [2,4-9]. In a 10-year study by Bengtson et al. [10] and in the study of Cunningham [11], after a 3-year follow-up period, 2.6% and 1.1% of primary augmentations, 4.7% and 2.3% of augmentation revision procedures, 4.9% and 2.5% of primary reconstruction procedures, and 3.0% and 1.5% of revision reconstruction procedures demonstrated implant malposition [10].

Numerous factors influence implant rotation [12]. A lack of connective tissue adhesion between the implant and the capsule can occur, prompting movement of the implant in the pocket [5]. Heitmann et al. [13], in their report, found that six of 132 patients experienced implant rotation because the surgical pocket was too large. Other factors that contribute to implant rotation include capsular fluid collection and prosthetic massage [14,15]. In addition, double capsules and dynamic action of the muscle, which may induce the implant to move laterally or vertically, also have been reported as possible factors contributing to implant rotation [16,17].

Meanwhile, anaplastic large cell lymphoma (ALCL) has become a major issue in breast augmentation and breast reconstruction surgery. In a recent study, it was reported that ALCL is mainly diagnosed in textured implants and may have some association with the textured surface, although the data remain too preliminary to draw any definitive conclusions [18]. A cumulative review of patients' status showed that ALCL is more prevalent in implants with aggressive texturing [19]. One of the most important reasons for using textured surfaces in anatomically shaped implants is to promote adhesion between the surface of the textured implant and the capsule, which is expected to prevent implant migration [20]. It is well-known that aggressively textured implant surfaces are associated with an increased risk of ALCL, meaning that the relationship between adhesion and ALCL is emerging as a noteworthy research topic.

However, to date, it is not known which parts of the implant are adhesively bonded to tissue, and which factors influence adhesion. In this study, we analyzed the adhesion pattern of tissue expanders, which are not permanent implants, but have similar texturing and are placed in the human body for several months. Thus, we sought to identify the patterns of adhesion development between textured surfaces and tissues in the human body, and which factors promote or degrade it.

METHODS

From September 2016 to November 2017, 79 tissue expanders were implanted in 75 patients at our facility as part of two-staged implant-based breast reconstruction procedures. Of these 79 tissue expanders, 14 were Mentor CPX4 expanders (Mentor Worldwide LLC, Irvine, CA, USA) and 65 were Natrelle expanders (Allergan plc, Dublin, Ireland). After mastectomy, acellular dermal matrix (ADM) was sutured along the inframammary fold and laterally to the chest wall. The superior part of the ADM was then sutured to the inferior border of the pectoralis major muscle to create a subpectoralis major muscle and ADM pocket. In 48 cases, CGCryoDerm (Daewoong Bio Inc., Seoul, Korea) was used, while MegaDerm (L&C Bio Co., Ltd., Seongnam, Korea) was used in 22 cases and DermACELL (LifeNet Health, Virginia Beach, VA, USA) was used in 10 cases. The size of the ADM varied from 4 × 16 to 6 × 17 cm. In all cases, the tissue expanders were inserted into the subpectoralis major muscle and acellular ADM pocket. Postoperatively, all patients used a Surgi-Bra (Golda Inc., Cleveland, OH, USA) continuously for 4 weeks and a nonwired sports bra for 2 weeks thereafter.

The presence of an adhesion area between the tissue expander and the capsule was investigated during the second operation, in which the tissue expander was exchanged for a permanent implant. We used nearly the same incision site as the previous mastectomy incision. A designated single operator checked for areas of adhesion between the tissue expander and capsule by dissection with a finger before removing the tissue expander in all cases. We calculated the interval from the immediate tissue expander reconstruction to the time of the second operation. We divided the tissue expanders into four areas: anterior-cephalic, anterior-caudal, posterior-cephalic, and posterior-caudal (Fig. 1), and categorized the patterns of adhesion into three types according to the number of areas that adhered. Total adhesion meant that adhesion had occurred in four areas, partial adhesion meant that adhesion had occurred in one to three areas, and non-adhesion meant that adhesion did not occur in any area. We investigated patient characteristics, such as tissue expander type, age, body mass index, total inflation amount, chemotherapy status, radiotherapy status, total Hemovac drainage amount, Hemovac drain removal date, the brand of ADM, and mastectomy type (total, skin-sparing, or nipple-sparing). Outcomes (number of attachments) were first analyzed with univariate analysis and then with multivariate logistic regression to quantify the impact of relevant patient demographics. Odds ratios were estimated using a logistic regression model. Patient demographics were summarized as means with standard deviations for continuous variables and as median values with interquartile

ranges (Q1, Q3) for continuous variables that did not show a normal distribution. Chi-square analysis or the Fisher exact test was used for categorical variables, the Mann-Whitney U test was used for continuous variables that did not show a normal distri-

bution, and the independent t-test was used for other continuous variables.

All statistical analyses were conducted using R version 3.4.4 (The R Foundation for Statistical Computing, Vienna, Austria).

RESULTS

There were 18 cases of total adhesion, 46 cases of partial adhesion, and 15 cases of non-adhesion. In terms of adhesion location, there were 58 cases of anterior-cephalic adhesion (58/64, 90.6%), 43 cases of anterior-caudal adhesion (43/64, 67.2%), 41 cases of posterior-cephalic adhesion (41/64, 64.1%), and 26 cases of posterior-caudal adhesion (26/64, 40.6%). There were three cases of minimal infections, which were addressed using oral antibiotics. All three of those infection cases involved a Natrelle expander and showed partial adhesion (posterior-caudal adhesion in one case and anterior-caudal adhesion in two cases).

The interval between the first operation and the second operation, in which the tissue expander was exchanged with a permanent breast implant, ranged from 4 months to 22 months (mean, 12 months). In the second operation, in which the expander was converted to a permanent implant, we used an Allergan Natrelle Style 410 implant in 55 cases and a Mentor CPG

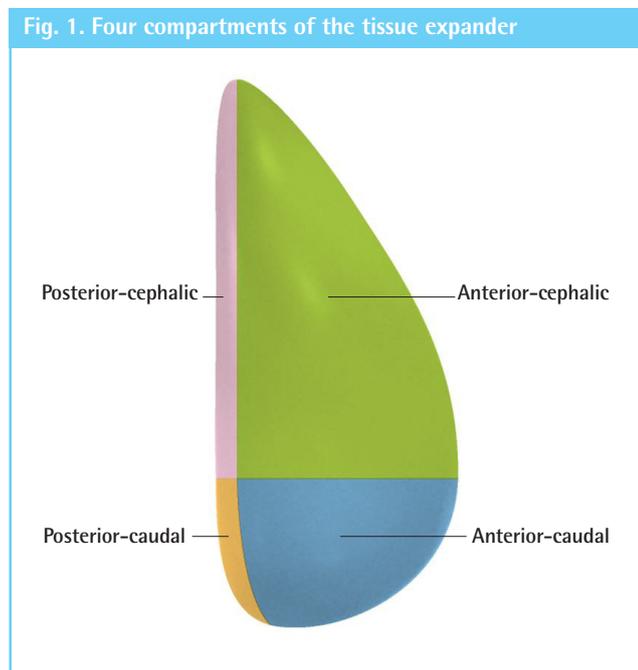


Table 1. Demographic and clinical information of the patients

Variable	Total (n = 79)	No. of attachments		P-value
		0 (n = 15)	≥ 1 (n = 64)	
Tissue expander brand				< 0.001
Allergan	65 (82.28)	3 (20.00)	62 (96.88)	
Mentor	14 (17.72)	12 (80.00)	2 (3.12)	
Age (yr)	45.54 ± 9.57	44.07 ± 8.55	45.89 ± 9.83	0.510
BMI (kg/m ²)	21.80 (20.23–24.34)	20.68 (19.73–23.38)	22.18 (20.39–24.41)	0.100
Total inflation	470.00 (360.00–520.00)	460.00 (370.00–510.00)	470.00 (357.50–520.00)	0.970
Chemotherapy				0.753
No	34 (43.04)	7 (46.67)	27 (42.19)	
Yes	45 (56.96)	8 (53.33)	37 (57.81)	
Radiotherapy				0.331
No	20 (25.32)	2 (13.33)	18 (28.12)	
Yes	59 (74.68)	13 (86.67)	46 (71.88)	
HV removal date	19.00 (16.50–21.00)	20.00 (18.00–21.00)	19.00 (15.00–21.00)	0.311
Total HV drainage amount	2,217.00 (2,079.00–2,384.50)	2,365.00 (2,246.50–2,437.50)	2,195.00 (2,055.25–2,357.75)	0.015
ADM type				> 0.999
CGcryoDerm	47 (59.49)	9 (60.00)	38 (59.38)	
MegaDerm	22 (27.85)	4 (26.67)	18 (28.12)	
DermACELL	10 (12.66)	2 (13.33)	8 (12.5)	
Interval from first to second operation	153.00 (119.50–284.50)	228.00 (167.50–259.00)	137.50 (117.00–296.75)	0.111
Mastectomy type				0.727
Total mastectomy	24 (30.38)	4 (26.67)	20 (31.25)	
Skin-sparing mastectomy	11 (13.92)	1 (6.67)	10 (15.62)	
Nipple-sparing mastectomy	44 (55.7)	10 (66.67)	34 (53.12)	

Values are presented as number (%), mean ± SD, or median (interquartile range). BMI, body mass index; HV, Hemovac; ADM, acellular dermal matrix.

implant in 24 cases. There was one case of rotation, which was confirmed to show 90° clockwise rotation at 12 months. This involved a Mentor CPGTM333 430-mL implant (tall height; high projection; width, 12 cm; height, 12.5 cm; projection, 6 cm) that showed non-adhesion, which was resolved by capsulectomy and change of the implant to Allergan Natrelle Style 410.

The patients' demographic and clinical information is summarized in Table 1. Seventeen cases of total adhesion were noted with the Natrelle expander, while only one case of total adhesion was noted with the Mentor CPX4 expander. The brand of tissue expander and the total Hemovac drainage amount showed statistically significant relationships with the number of attachments ($P < 0.001$ and $P = 0.015$, respectively). No statistical significance was found for the relationships of the mastectomy type and interval between the first and second operations with the number of attachments. When the cases were stratified by the tissue expander brand, non-adhesion (attachment number = 0) was observed in 20% of cases using an Allergan Natrelle expander and in 80% of cases using a Mentor CPX4. The Natrelle expander and Mentor CPX4 showed a significant difference in the number of cases with at least one attachment (96.9% and 3.1%, respectively) (Table 1).

Additionally, in the multivariable logistic results, only the tis-

sue expander brand showed a statistically significant relationship with adhesion ($P < 0.001$). Therefore, the tissue expander brand was found to be the variable with the most influence on the number of attachments (Table 2).

DISCUSSION

In this study, we divided the areas of the tissue expanders into four categories based on anatomical location, and investigated which areas most frequently adhered to the surrounding tissues. We also studied patient-related factors that affected adhesion and attempted to determine how much adhesion was required to avoid malrotation of the implant.

In implant-based breast reconstruction and breast augmentation, breast implant adhesion refers to fibrosis with the capsule and implant. In a study by Smahel et al. [21], the authors stated that the soft tissue response was different depending on the texture of the implant used. For 6–8 months after implantation, at the microscopic level, the number of myofibroblasts increased as compared with the smooth side and collagen deposition was more progressive in the “bridges” than at the base of villous processes. After 8 months, the basal layer of the capsule was fibrotic. As a result, at the microscopic level, cavities were present in the

Table 2. Logistic regression analysis

Variable	Univariable logistic regression		Multivariable logistic regression	
	OR (95% CI)	P-value	OR (95% CI)	P-value
Tissue expander brand				
Allergan	Reference		Reference	
Mentor	0.008 (0.001–0.054)	< 0.001	0.009 (0.001–0.063)	< 0.001
Age	1.021 (0.961–1.083)	0.505		
BMI	1.242 (0.986–1.565)	0.066		
Total inflation	0.999 (0.994–1.005)	0.845		
Chemotherapy				
No	Reference			
Yes	1.199 (0.388–3.709)	0.753		
Radiotherapy				
No	Reference			
Yes	0.393 (0.081–1.919)	0.248		
Hemovac removal date	0.927 (0.831–1.035)	0.179		
Total Hemovac drainage amount	0.998 (0.996–1.000)	0.036	0.999 (0.996–1.003)	0.710
ADM type				
CGcryoDerm	Reference			
MegaDerm	1.066 (0.289–3.928)	0.924		
DermACELL	0.947 (0.171–5.245)	0.951		
Interval from first to second operation	0.999 (0.995–1.004)	0.763		
Mastectomy type				
Total mastectomy	Reference			
Skin-sparing mastectomy	2.000 (0.197–20.325)	0.558		
Nipple-sparing mastectomy	0.680 (0.188–2.456)	0.556		

Outcome of logistic regression analysis: attachment number (0 vs. ≥ 1).
OR, odds ratio; CI, confidence interval; BMI, body mass index.

textured implant surface. They called these cavities “bridges.” Fibrosis in these bridges led to better adhesion between the capsule and implant. Based on those results, it could be predicted that more aggressive implant texturing induces more adhesion.

In the current study, we found that the extent to which capsule and tissue expander adhesion was induced depended on the brand of tissue expander used. The Natrelle expander demonstrated more instances of adhesion than did the Mentor CPX4 expander. Due to the differences in the manufacturing process, the Natrelle expander has a more aggressively textured surface than Mentor CPX4, as shown by microscopy [22]. This means that the texture of the implant is associated with adhesion. In a study by Montemurro et al. [12], the authors stated their belief that good adhesive properties contributed to a lower rotation rate. Furthermore, Danino et al. [23] demonstrated that a sufficient texture contributed to a more stable adhesive effect involving the surrounding capsule. A prospective study of electron microscopic findings in adhesion and non-adhesion areas could yield insights into breast implant adhesion [21].

In our study, the anterior part of the expander was the area that was most adhesive to the capsule, and the posterior-caudal part was the area that was most non-adhesive to the capsule. The anterior part of the expander is the area where the least fluid collects, because it is a non-dependent area, while the posterior-caudal part is the area where the most fluid collects because it is the dependent area of the pocket. Seroma was found to have an effect on adhesion between the implant and capsule [14], suggesting that differences in fluid collection may be a contributing factor to this finding.

There were no instances of malrotation in cases of tissue expander insertion in our study, although there was one case of malrotation that showed non-adhesion after the second operation. Even in the 15 expanders with non-adhesion, malrotation did not occur. A reason for this may be that the observer missed slight malrotation. In addition, it is presumed that the frictional force of the expander itself prevented malrotation, even without adhesion.

Partial adhesion may be a common phenomenon after breast tissue expander insertion, and it is possible that it could occur at a similar rate in cases of breast augmentation or permanent implant placement in breast reconstruction. Although our study had a relatively short duration of observation, and did not observe the actual adhesion of permanent implants with capsules, we predict that even partial adhesion suffices to prevent expander rotation. Because of this limitation, in a further study we will be able to strengthen our argument by incorporating a longer period of observation and visual confirmation of the adhesion of permanent implants with capsules.

In the study of Maxwell et al. [8], appropriate adhesion of the implant to the surrounding tissue was found to be essential for maximizing the benefits of the procedure and securing a low rate of rotation. Maxwell et al. urged surgeons to ensure precise pocket dissection, with the drain inserted into a precisely fitting implant pocket and bloodless sharp dissection (with loose connective tissue left on the ribs to reduce bleeding), and to avoid massage and vigorous displacement exercises for up to 3 months, as those steps may promote tissue adhesion and prevent implant rotation.

However, our study suggests that aggressive texturing is a more important factor than those described by Maxwell et al. The aggressiveness of the texture of the implant was found to be the most important factor associated with implant adhesion in this study. This result is noteworthy, since ALCL also occurs more frequently in aggressively textured implants. As the relationship between ALCL and textured implants is being actively researched, it is expected that further studies of the relationship between ALCL and the extent of adhesion will be of interest and shed light onto basic aspects of ALCL.

In this study, we evaluated the location of adhesion after tissue expander insertion. Our data indicated that the texture of the implant was a significant factor contributing to the success of adhesion. Partial adhesion is a common phenomenon in expander-based breast reconstruction. Nevertheless, partial adhesion was found to be sufficient to prevent unwanted rotation of the implant in this study. Because the anterior-cephalic surface showed the highest adhesion rate, the anterior portion of the implant may have the highest likelihood of successful adhesion to the capsule.

NOTES

Conflict of interest

No potential conflict of interest relevant to this article was reported.

Ethical approval

The study was and performed in accordance with the principles of the Declaration of Helsinki. Written informed consents were obtained.

Author contribution

Conceptualization: Song SY, Park KH. Data curation: Park KH, Lim YM. Formal analysis: Lim YM. Methodology: Lee DW. Project administration: Lew DH, Roh TS. Writing - original draft: Park KH. Writing - review & editing: Lim YM. Approval of final manuscript : all authors.

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