

# A Prospective, Randomized Comparison of Clinical Outcomes with Different Processing Techniques in Autologous Fat Grafting

Summer E. Hanson, M.D.,  
Ph.D.

Patrick B. Garvey, M.D.

Edward I. Chang, M.D.

Gregory P. Reece, M.D.

Jun Liu, Ph.D.

Donald P. Baumann, M.D.

Charles E. Butler, M.D.

Chicago, Ill.; and Houston, Texas



**Background:** Autologous fat grafting is a useful tool in breast reconstruction. The authors have previously demonstrated a difference in the rate of processing adipose grafts in a randomized time and motion clinical trial. The purpose of this study was to compare clinical outcomes in commonly used grafting systems.

**Methods:** Three methods to prepare adipose grafts were compared: a passive washing filtration system (Puregraft system), an active washing filtration system (Revolve system), and centrifugation (Coleman technique). Postoperative complications, rates of fat necrosis, revision procedures, and additional imaging were recorded.

**Results:** Forty-six patients were included in the prospective, randomized study (15 active filtration, 15 passive filtration, and 16 centrifugation). Their mean age was 54 years and mean body mass index was 28.6 kg/m<sup>2</sup>. The mean length of follow-up was 16.9 ± 4 months. The overall complication rate was 12.1 percent. The probability of fat necrosis was no different between the groups (active filtration, 15 percent versus passive filtration, 14.3 percent] versus centrifugation, 8 percent;  $p = 0.72$ ). Fat necrosis was highest in patients with breast conservation before grafting (60 percent;  $p = 0.011$ ). There was no significant difference in contour irregularity (active filtration, 40 percent versus passive filtration, 38 percent versus centrifugation, 36 percent;  $p = 0.96$ ) or additional grafting (active filtration, 40 percent versus passive filtration, 24 percent versus centrifugation, 32 percent;  $p = 0.34$ ).

**Conclusions:** This is the first prospective, randomized study to compare clinical outcomes of adipose graft preparation. There was no significant difference in early complications, fat necrosis, or rates of additional grafting among the study groups. There was significantly higher risk of fat necrosis in patients with previous breast conservation treatment regardless of processing technique. (*Plast. Reconstr. Surg.* 150: 955, 2022.)

**CLINICAL QUESTION/LEVEL OF EVIDENCE:** Therapeutic, II.

**A**utologous fat grafting is a useful tool in cosmetic and reconstructive surgery. In recent years, the indications in soft-tissue reconstruction have expanded beyond volume loss and contour deformity<sup>1-4</sup> to include radiation fibrosis, peripheral neuropathy, complex pain, and lymphedema.<sup>5,6</sup> The underlying

mechanisms of these benefits are not fully understood, but the effects of purified adipose tissue are just as likely to be paracrine as physical. With the increase in procedural volume<sup>7</sup> comes increasing interest in optimizing the process, from preparing a more robust graft to reducing operative time and cost. There are three key steps in grafting—harvest, processing,

From the Department of Surgery, University of Chicago Medicine and Biological Sciences; and Department of Plastic Surgery, The University of Texas M. D. Anderson Cancer Center.

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and delivery—each of which has several variables that may affect the efficiency and efficacy of the procedure.

The primary objective in processing the lipoaspirate before injection is to remove impurities that have resulted from tissue harvest, such as cellular debris, lipid, or blood. There are several described techniques to prepare a graft with no one method being shown to be superior to others.<sup>8–10</sup> Members of the American Society of Plastic Surgeons were previously surveyed and reported preferred processing techniques including centrifugation (34 percent), filtration (34 percent), and washing or rinsing (28 percent).<sup>11</sup> Since that survey was conducted, device-based processing systems have gained popularity for their streamlined, closed handling.

The Revolve system (LifeCell Corporation, Branchburg, N.J.)<sup>12</sup> is an active filtration system that uses mechanical suction to harvest and filter the lipoaspirate through a closed loop canister. Standard liposuction can be performed and resulting debris is filtered through a 75- $\mu$ m mesh sieve. The Puregraft 250 system (Cytori Therapeutics, San Diego, Calif.)<sup>13</sup> is a passive filtration system that similarly uses a lactated Ringer washing base step. Lipoaspirate can be harvested through handheld syringe or machine-assisted suction. The collection bag in this system is a double-layer mesh filter with the inner mesh being 50  $\mu$ m and the outer mesh being 1000  $\mu$ m. Graft preparation by centrifugation is the most commonly studied in the literature and has been varied by both time and speed. The method used in this study is similar to that described by Coleman<sup>1</sup> of 3000 revolutions per minute for 3 minutes. With either of the filtration devices, intact lobules of graft are preserved and smaller debris is filtered to the waste canister while centrifugation effectively separates the components by density.

We previously performed a prospective, randomized time and motion study comparing the operative outcomes of three commonly used processing techniques.<sup>14</sup> There was significant difference in the rate of processing the tissue graft in each system, favoring the active filtration device. The purpose of the current study is to compare short-term and 1-year clinical outcomes of the three commonly used systems for graft preparation. Our hypothesis was that preparation with centrifugation will have higher rates of fat necrosis and additional grafting compared with the devices that include washing and filtration.

## PATIENTS AND METHODS

### Study Design

The authors conducted a randomized, prospective parent study to compare outcomes in processing techniques for fat grafting.<sup>14</sup> The protocol was approved by the authors' institutional review board (P2015-0006) as well as the Clinical Oncology Research system (CORG) and registered with the U.S. National Library of Medicine at ClinicalTrials.gov (no. NCT02677012). The parent study was designed, and therefore powered, to identify a difference in rate of processing the adipose graft among the three study arms.<sup>15</sup> Overall, 46 participants were included: 15 per arm with one who failed final screening. Exclusion criteria were age younger than 18 years, inability to give consent, active chemotherapy or cancer treatment, pregnancy, body mass index less than 18, or lacking an anticipated minimum volume of 100 mm of lipoaspirate.

Consecutive patients were stratified by a core group of participating surgeons and randomized by technique 1:1:1 for each arm. Once the eligible participant was registered with the CORG system, CORG randomly assigned the patient to one of three fat grafting techniques using one of five surgeon blocks as the allocation stratum.

Patient information collected included age, body mass index, previous medical and surgical history, oncologic history including tumor characteristics, surgery, chemotherapy or irradiation, and reconstructive characteristics. Follow-up information included early complications such as seroma or hematoma diagnosed on examination or ultrasound but not necessarily requiring drainage or cellulitis of the donor or recipient site treated with antibiotics within 30 days after grafting. Long-term outcomes included any palpable mass appreciated on physical examination persistent for more than 3 months, contour irregularity noted by physical examination and persistent for more than 3 months, fat necrosis identified by imaging (mammography or ultrasound), other radiographic abnormality such as calcifications or oil cyst, need for tissue biopsy in the grafted breast, or development of cancer in the grafted breast. Contour irregularity was a subjective assessment made by the plastic surgeon at the follow-up visits. It is proposed by Delay and colleagues<sup>2</sup> that "steady state" volume and contour effects of grafting are achieved by approximately 3 months and therefore "persistent" contour irregularity referred to a difference in the contour of the area grafted that was present beyond 3 months when

we would expect soft-tissue edema to resolve postoperatively. This was most commonly a contracted scar or upper pole step-off deformity that did not fully improve with grafting. In reference to the donor site, this refers to a contour irregularity from the liposuction that was present more than 3 months postoperatively, again when we would expect to see the resolution of tissue edema from the procedure. Additional grafting procedures that occurred in the follow-up period were recorded.

**Statistical Methods**

Descriptive statistics, such as means and standard deviations, were used to summarize continuous variables. Frequency counts and percentages were used for categorical variables. One-way analysis of variance was applied to evaluate the difference of age and body mass index among the three groups. Kruskal-Wallis tests were used to compare the tumor size and volume injected to breasts. Post hoc comparisons were performed to identify individual groups. The Tukey studentized range (honestly significant difference) test was used to adjust multiple comparisons. The chi-square or Fisher exact test was employed to assess the associations between oncologic characteristics, procedures, autologous fat grafting procedure, and postoperative complications and the techniques. Univariate logistic regression model was used to estimate the effect of techniques and clinical factors on probability of fat necrosis. The Hosmer and Lemeshow test was performed to examine the goodness of fit for the logistic regression model. The area under the receiver operating characteristic curve was used to evaluate the predictive probability. All significance tests were two-sided and *p* value less than

5 percent was considered statistically significant. All data analyses were performed using SAS 9.4 software (SAS Institute, Cary, N.C.).

**RESULTS**

**Patient Characteristics**

A total of 46 patients were enrolled in the parent study.<sup>14</sup> Fifteen were randomized to active filtration, 15 were randomized to passive filtration, and 16 were randomized to centrifugation, with one failing the initial screen. There were no statistical differences in patient characteristics among the three groups (Table 1). All patients were women who had undergone fat grafting for reconstruction related to breast cancer. Their mean age was 54 ± 10 years and their mean body mass index was 28.6 ± 4.1 kg/m<sup>2</sup>. There were similar rates of tobacco use and comorbidities. All patients were treated previously for breast cancer. The most common diagnosis of participants was ductal carcinoma in situ (40.9 percent), with an average overall tumor size of 1.9 ± 2.1 cm. There was no difference in histology, stage, or tumor location among the processing groups (Table 2).

**Surgical Characteristics**

Reconstructive characteristics are summarized in Table 3 and analysis was performed on the breast level. The majority of patients had previously undergone mastectomy with two-stage reconstruction. Just over half had definitive autologous reconstruction (53 percent) and only five patients had breast-conserving therapy including whole breast irradiation (7.6 percent). Regarding the fat grafting procedures (Table 4), operations

**Table 1. Patient Characteristics by Technique and Overall Intention-to-Treat Population\***

Variable	All (n = 46)	Active Filtration System (n = 15)	Passive Filtration System (n = 15)	Centrifugation (n = 16)	<i>p</i>
Age at AFG, yrs	54.0 ± 10.10	55.5 ± 10.2	52.7 ± 11	53.9 ± 9.6	0.79
BMI, kg/m <sup>2</sup>	28.6 ± 4.1	30.2 ± 4.3	28.4 ± 3.3	27.2 ± 4.3	0.03†
Race					0.45
White	33 (71.7)	10 (66.7)	11 (73.3)	12 (75)	
Black	6 (13)	4 (26.7)	1 (6.7)	1 (6.3)	
Hispanic	7 (15.2)	1 (6.7)	3 (20)	3 (18.8)	
Tobacco use					0.21
Never	36 (78.3)	10 (66.7)	12 (80)	14 (87.5)	
Current	1 (2.2)	0 (0)	0 (0)	1 (6.3)	
Former	9 (19.6)	5 (33.3)	3 (20)	1 (6.3)	
Comorbidity	27 (58.7)	11 (73.3)	9 (60)	7 (43.8)	0.25
DM	3 (6.5)	2 (13.3)	1 (6.7)	0 (0)	0.30
HTN	15 (32.6)	7 (46.7)	6 (40)	2 (12.5)	0.08
Cardiac	13 (28.3)	7 (46.7)	5 (33.3)	1 (6.3)	0.03†
Pulmonary	2 (4.3)	1 (6.7)	0 (0)	1 (6.3)	1.000
Digestive	11 (23.9)	4 (26.7)	3 (20)	4 (25)	1.000

AFG, autologous fat grafting; BMI, body mass index; DM, diabetes mellitus; HTN, hypertension.

\*Values are expressed as mean ± SD or *n* (%).

†Statistically significant.

**Table 2. Summary of Oncologic Characteristics per Breast by Technique\***

Variable	All (n = 66)	Active Filtration System (n = 20)	Passive Filtration System (n = 21)	Centrifugation (n = 25)	p
Histology					0.47
IDC	23 (34.8)	5 (25)	8 (38.1)	10 (40)	
DCIS	27 (40.9)	12 (60)	7 (33.3)	8 (32)	
ILC	4 (6.1)	2 (10)	1 (4.8)	1 (4)	
Mixed	7 (10.6)	1 (5)	3 (14.3)	3 (12)	
LCIS	4 (6.1)	0 (0)	1 (4.8)	3 (12)	
Other	1 (1.5)	0 (0)	1 (4.8)	0 (0)	
Stage					0.34
0	30 (45.5)	5 (25)	12 (57.1)	13 (52)	
1	8 (12.1)	2 (10)	2 (9.5)	4 (16)	
2	16 (24.2)	8 (40)	4 (19)	4 (16)	
3	12 (18.2)	5 (25)	3 (14.3)	4 (16)	
Location					0.82
UOQ	22 (33.3)	6 (30)	6 (28.6)	10 (40)	
UIQ	20 (30.3)	5 (25)	7 (33.3)	8 (32)	
LOQ	5 (7.6)	2 (10)	2 (9.5)	1 (4)	
LIQ	2 (3)	1 (5)	0 (0)	1 (4)	
Multicentric	3 (4.5)	2 (10)	0 (0)	1 (4)	
Upper central	3 (4.5)	0 (0)	1 (4.8)	2 (8)	
Lower central	11 (16.7)	4 (20)	5 (23.8)	2 (8)	
Chemotherapy	31 (47)	12 (60)	10 (47.6)	9 (36)	0.28
Radiation	20 (30.3)	9 (45)	4 (19)	7 (28)	0.19
BRCA	12 (18.2)	6 (30)	0 (0)	6 (24)	0.03†
Tumor size, cm	1.5 (0–8.9)	2 (0–8.9)	1.5 (0–7.5)	0.8 (0–8)	0.34

AFG, autologous fat grafting; DCIS, ductal carcinoma in situ; IDC, invasive ductal carcinoma; ILC, invasive lobular carcinoma; LCIS, lobular carcinoma in situ; LIQ, lower inner quadrant; LOQ, lower outer quadrant; UIQ, upper inner quadrant; UOQ, upper outer quadrant.

\*Values are expressed as n (%) or median (range).

†Statistically significant.

**Table 3. Summary of Oncologic Procedure per Breast by Technique\***

Characteristic	All (n = 66)	Active Filtration System (n = 20)	Passive Filtration System (n = 21)	Centrifugation (n = 25)	p
Side					0.71
Left	46 (69.7)	15 (75)	15 (71.4)	16 (64)	
Right	20 (30.3)	5 (25)	6 (28.6)	9 (36)	
Breast conservation	5 (7.6)	2 (10)	1 (4.8)	2 (8)	0.86
Mastectomy	61 (92.4)	18 (90)	20 (95.2)	23 (92)	0.86
Prophylactic	22 (33.3)	6 (30)	6 (28.6)	10 (40)	0.67
Reconstruction					
Oncoplastic	5 (7.6)	2 (10)	1 (4.8)	2 (8)	0.86
TE/implant	25 (37.9)	5 (25)	7 (33.3)	13 (52)	0.16
ADM	24 (36.4)	5 (25)	6 (28.6)	13 (52)	0.12
Autologous	34 (51.5)	12 (60)	12 (57.1)	10 (40)	0.34

ADM, acellular dermal matrix; AFG, autologous fat grafting; DIEP, deep inferior epigastric perforator; TE, tissue expander; TRAM, transverse rectus abdominus myocutaneous.

\*Values are expressed as n (%).

were more commonly bilateral and performed with additional procedures. The average volume injected per breast was 89.7 ± 63.9 ml with slightly less in the centrifugation cohort (96.8 ml active filtration versus 99.8 ml passive filtration versus 75.7 ml centrifugation; *p* = 0.44). The majority of cases were performed with other procedures, commonly scar revision, contralateral mastopexy for symmetry, or tissue expander to implant exchange.

**Complications and Long-term Outcomes**

The overall complication rate was 12.1 percent, all of which was fat necrosis, with similar incidence

among the processing techniques (Table 5; 15 percent active filtration versus 14.3 percent passive filtration versus 8 percent centrifugation; *p* = 0.72). There was no occurrence of cellulitis, hematoma, or seroma within 30 days of grafting. There was no occurrence of locoregional recurrence, metastatic disease, or development of new cancer (invasive or in situ) during the study period. More than one-third of patients had persistent contour irregularity at the breast level (40 percent active filtration versus 38.1 percent passive filtration versus 36 percent centrifugation; *p* = 0.3) with statistically similar rates of additional fat grafting (20 percent active filtration versus 14.3 percent

**Table 4. Summary of Autologous Fat Grafting Procedure per Breast by Technique\***

Characteristic	All	Active Filtration System (n = 20)	Passive Filtration System (n = 21)	Centrifugation (n = 25)	p
Unilateral AFG†	26 (39.4)	10 (50)	9 (42.9)	7 (28)	0.30
Bilateral AFG	40 (60.6)	10 (50)	12 (57.1)	18 (72)	0.30
Previous AFG	17 (25.8)	2 (10)	6 (28.6)	9 (36)	0.13
Donor site					
Abdomen/flanks	52 (78.8)	16 (80)	19 (90.5)	17 (68)	0.17
Thigh	45 (68.2)	13 (65)	14 (66.7)	18 (72)	0.87
Buttocks	2 (3)	0 (0)	0 (0)	2 (8)	0.33
Additional procedures					
TE implant or exchange	12 (18.2)	3 (15)	0 (0)	9 (36)	0.004‡
Mastopexy	19 (28.8)	5 (25)	6 (28.6)	8 (32)	0.88
Scar revision	52 (78.8)	17 (85)	19 (90.5)	16 (64)	0.09
Other	20 (30.3)	6 (30)	9 (42.9)	5 (20)	0.24
AFG volume, ml					
Total procedure	112 (33–385)	109.5 (60–265)	114 (60–385)	112 (33–260)	0.79
Per breast	77 (17–385)	108 (17–215)	75 (30–385)	70 (25–150)	0.44

AFG, autologous fat grafting; TE, tissue expander.  
 \*Values are expressed as n (%) or median (range).  
 †Unilateral autologous fat grafting to the cancer-affected breast.  
 ‡Statistically significant.

**Table 5. Summary of Postoperative Complications per Breast by Technique\***

Complication	All (n = 66)	Active Filtration System (n = 20)	Passive Filtration System (n = 21)	Centrifugation (n = 25)	p
Any	8 (12.1)	3 (15)	3 (14.3)	2 (8)	0.72
Hematoma	0 (0)	0 (0)	0 (0)	0 (0)	—
Seroma	0 (0)	0 (0)	0 (0)	0 (0)	—
Cellulitis	0 (0)	0 (0)	0 (0)	0 (0)	—
Fat necrosis	8 (12.1)	3 (15)	3 (14.3)	2 (8)	0.72
Palpable mass	7 (10.6)	2 (10)	3 (14.3)	2 (8)	0.88
Contour change					
Breast	25 (37.9)	8 (40)	8 (38.1)	9 (36)	0.30
Donor site	1 (1.5)	1 (5)	0 (0)	0 (0)	0.35
Recurrence	0 (0)	0 (0)	0 (0)	0 (0)	—
Metastasis	0 (0)	0 (0)	0 (0)	0 (0)	—
New cancer	0 (0)	0 (0)	0 (0)	0 (0)	—
Additional AFG	15 (22.7)	4 (20)	3 (14.3)	8 (32)	0.35

AFG, autologous fat grafting.  
 \*Values are expressed as n (%).

**Table 6. Univariate Analysis of Probability of Fat Necrosis in All Processing Groups Combined\***

Variable	No Fat Necrosis (n = 58)	Fat Necrosis (n = 8)	p
Processing technique			0.715
System active filtration	17 (85)	3 (15)	
System passive filtration	18 (85.7)	3 (14.3)	
Centrifugation	23 (92)	2 (8)	
Chemotherapy	28 (90.3)	3 (9.7)	0.713
Radiation therapy	17 (85)	3 (15)	0.690
Oncologic surgery			0.011†
Breast conservation	2 (40)	3 (60)	
Mastectomy	56 (91.8)	5 (8.2)	
Prophylactic	21 (95.5)	1 (4.5)	0.252
Reconstruction			
Oncoplastic	2 (40)	3 (60)	0.020†
TE/implant	24 (96)	1 (4)	0.143
ADM	23 (95.8)	1 (4.2)	0.241
Autologous (any flap)	30 (88.2)	4 (11.8)	1.000
Mastopexy	14 (73.7)	5 (26.3)	0.038†
Total grafted volume, ml	111 (33–385)	130 (75–150)	0.694
Breast grafted volume, ml	71 (17–385)	87.5 (50–130)	0.330

ADM, acellular dermal matrix; TE, tissue expander.  
 \*Values are expressed as n (%) or median (range).  
 †Statistically significant.

passive filtration versus 32 percent centrifugation; *p* = 0.35). Univariate analysis of fat necrosis demonstrates significantly higher probability with

breast-conserving therapy including irradiation, oncoplastic surgery, or mastopexy irrespective of processing technique (Table 6).

## DISCUSSION

This is the first prospective, randomized comparison of two commonly used closed-system devices and centrifugation for preparation of lipoaspirate for autologous fat grafting. Despite different means of purifying the tissue graft, there was no statistical difference in rates of fat necrosis, residual contour deformity, or additional grafting. However, we demonstrate a higher probability of fat necrosis with breast-conserving therapy, oncoplastic reconstruction, or mastopexy regardless of processing technique.

The indications and procedural volume of fat grafting cases continue to increase, particularly in oncologic reconstruction<sup>16-18</sup>; however, there is no evidence to suggest one technique is clinically better than others.<sup>19</sup> This may be attributable in part to inconsistency in outcome measures, length of follow-up, and reporting. Particularly in breast reconstruction, average complication rates reported are 8.4 percent (95 percent CI, 7.6 to 9.1 percent), most commonly cyst or fat necrosis.<sup>20</sup> Although these outcomes are generally considered benign, any breast abnormality in a patient previously treated for breast cancer may cause stress and lead to additional imaging or biopsy.<sup>17,21-23</sup> Thus, our long-term research goal is to determine optimal techniques and establish best practices to improve outcomes of autologous fat grafting in breast reconstruction.

Rates of fat necrosis in the setting of breast-conserving therapy with more than 6 years follow-up are comparable, with or without fat grafting.<sup>17</sup> This similarity is likely attributable to the changes that occur with any procedure that alters the parenchyma as in reduction mammoplasty, mastopexy, or lumpectomy.<sup>24,25</sup> In the current study, our data demonstrate higher probability of fat necrosis with breast-conserving therapy and parenchymal reshaping procedures including oncoplastic reconstruction or revision mastopexy. Thus, in counseling patients, we know breast conservation has a higher likelihood of fat necrosis of the breast related to the combination of lumpectomy surgery and irradiation. It is important to discuss any fat necrosis before grafting that may have resulted from oncologic treatment.

There was no difference in rates of fat necrosis in the setting of irradiation in the current study; however, these data should be scrutinized more thoroughly, particularly in participants who underwent mastectomy. There were no participants in any cohort who had implant-based reconstruction followed by postmastectomy irradiation, nor were

there patients with autologous breast reconstruction who then underwent irradiation. This is a reflection of the authors' overall practice favoring autologous reconstruction, especially when radiation treatment is indicated. Although irradiation did not correlate with increased risk of fat necrosis in subsequent fat grafting, all patients in this study who had mastectomy followed by postmastectomy irradiation underwent autologous breast reconstruction. This means that after mastectomy, adipose graft was delivered to a vascularized recipient bed in the autologous tissue flap. Any fat grafting that occurred in the setting of implant-based reconstruction was to nonirradiated mastectomy skin flaps, pointing to the value of a well-vascularized recipient bed in mitigating ischemia in fat grafting outcomes. We can conclude, however, that alloplastic reconstruction including the use of acellular dermal matrix does not carry additional risk of fat necrosis in the absence of breast irradiation regardless of processing technique.

The majority of retrospective studies available include centrifugation or decanting as the processing technique.<sup>2,4,18,20,26-28</sup> As the popularity of autologous fat grafting expands, numerous methods of harvesting, processing, and reinjecting fat have evolved, most notably devices or systems to streamline the process.<sup>29</sup> Few studies have evaluated the outcomes specific to tissue processing technique. A recent retrospective review compared clinical outcomes in fat grafting cases to the breast among centrifugation, Telfa rolling (in which the tissue is gently rolled along a gauze sponge to remove fluid), and Revolve (or the active filtration system employed in this study).<sup>30</sup> The study included 267 cases with 12-month follow-up and found overall complication rates were highest in the centrifugation group, at 25.6 percent, compared with 4.5 percent with Telfa and 10.9 percent with the Revolve filtration system ( $p = 0.001$ ); rates of fat necrosis were similar and oil cysts accounted for the difference. The average volume of fat injected was significantly higher in the Revolve group (160 ml) compared with Telfa (120 ml) and centrifugation (70 ml). The authors' multivariate analysis found more "adverse events" with higher graft volume and centrifugation technique. Although our data show less volume of graft per breast with the centrifugation group (75 ml) compared with active filtration (99.5 ml) and passive filtration (96 ml), this did not achieve statistical significance. Overall, the volume of graft and processing technique did not have an effect on rates of fat necrosis in our prospective study.

Although our study was prospective with randomized study cohorts, the primary limitation is the sample size. The source time and motion study was powered to demonstrate a statistical difference in techniques based on time and motion methodology and clearly favored the active filtration system.<sup>14</sup> Given that reported rates of fat necrosis are low in autologous fat grafting, one would need to enroll nearly 1000 patients to achieve adequate statistical power to demonstrate a significant difference. Although the focus of this study was objective clinical outcomes, another limitation may be the lack of a validated patient-reported outcome survey. The authors believed that because there are no patient-reported outcomes specific to fat grafting, the BREAST-Q elements would be confounded by the fact that the majority of the participants in this study had grafting included with additional reconstructive procedures such as scar revision or tissue expander to implant exchange. Finally, we did not include specific volumetric analysis in this study. Graft retention is attributed to graft purity, recipient site preparation, optimal delivery technique, and ischemia.<sup>31-34</sup> Furthermore, graft retention is reported as ranging from 20 to 90 percent<sup>8</sup>; however, there are no reliable methods to quantify graft volume over time and most studies lack adequate follow-up to demonstrate long-term volume changes. For this reason, the authors included clinically relevant outcomes of complications and additional grafting. Optimal detection of graft volume and breast shape over time are under investigation at our institution.

## CONCLUSIONS

This is the first randomized prospective study of autologous fat grafting for oncologic breast reconstruction to compare techniques to process lipoaspirate. The authors found no statistical difference in complications or fat necrosis among centrifugation, an active filtration device (Revolve), or a passive filtration system (Puregraft 250). Instead, the data demonstrate a significantly higher probability of fat necrosis with breast-conserving therapy, including irradiation, oncoplastic surgery, or revision mastopexy.

**Summer E. Hanson, M.D., Ph.D.**

Plastic and Reconstructive Surgery  
University of Chicago Medicine and Biological Sciences  
5841 S. Maryland Avenue, Room J641  
Chicago, Ill. 60637  
sehanson@bsd.uchicago.edu  
@DrSummerHanson

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## REFERENCES

1. Coleman SR. Structural fat grafting. *Aesthet Surg J.* 1998;18:386, 388.
2. Delay E, Garson S, Tousson G, Sinna R. Fat injection to the breast: Technique, results, and indications based on 880 procedures over 10 years. *Aesthet Surg J.* 2009;29:360-376.
3. Gutowski KA; ASPS Fat Graft Task Force. Current applications and safety of autologous fat grafts: A report of the ASPS fat graft task force. *Plast Reconstr Surg.* 2009;124:272-280.
4. Myckatyn TM, Wagner IJ, Mehrara BJ, et al. Cancer risk after fat transfer: A multicenter case-cohort study. *Plast Reconstr Surg.* 2017;139:11-18.
5. Caviggioli F, Maione L, Forcellini D, Klinger F, Klinger M. Autologous fat graft in postmastectomy pain syndrome. *Plast Reconstr Surg.* 2011;128:349-352.
6. Caviggioli F, Maione L, Klinger F, Lisa A, Klinger M. Autologous fat grafting reduces pain in irradiated breast: A review of our experience. *Stem Cells Int.* 2016;2016:2527349.
7. American Society of Plastic Surgeons. 2016 Plastic Surgery Statistics Report: ASPS National Clearinghouse of Plastic Surgery Procedural Statistics. Arlington Heights, Ill.: American Society of Plastic Surgeons; 2017.
8. Gir P, Brown SA, Oni G, Kashefi N, Mojallal A, Rohrich RJ. Fat grafting: Evidence-based review on autologous fat harvesting, processing, reinjection, and storage. *Plast Reconstr Surg.* 2012;130:249-258.
9. Strong AL, Cederna PS, Rubin JP, Coleman SR, Levi B. The current state of fat grafting: A review of harvesting, processing, and injection techniques. *Plast Reconstr Surg.* 2015;136:897-912.
10. Kaufman MR, Bradley JP, Dickinson B, et al. Autologous fat transfer national consensus survey: Trends in techniques for harvest, preparation, and application, and perception of short- and long-term results. *Plast Reconstr Surg.* 2007;119:323-331.
11. Kling RE, Mehrara BJ, Pusic AL, et al. Trends in autologous fat grafting to the breast: A national survey of the American Society of Plastic Surgeons. *Plast Reconstr Surg.* 2013;132:35-46.
12. Allergan. Revolve Advanced Adipose System: Instructions for Use. Branchburg, NJ: Allergan; 2013.
13. Cytori Therapeutics. Puregraft 250 System: Instructions for Use. San Diego, Calif: Cytori; 2010.
14. Hanson SE, Garvey PB, Chang EI, et al. A randomized prospective time and motion comparison of techniques to process autologous fat grafts. *Plast Reconstr Surg.* 2021;147:1035-1044.
15. Hanson SE, Garvey PB, Chang EI, Reece G, Liu J, Butler CE. A prospective pilot study comparing rate of processing techniques in autologous fat grafting. *Aesthet Surg J.* 2019;39:331-337.
16. Karmali RJ, Hanson SE, Nguyen AT, Skoracki RJ, Hanasono MM. Outcomes following autologous fat grafting for oncologic head and neck reconstruction. *Plast Reconstr Surg.* 2018;142:771-780.

17. Hanson SE, Kapur SK, Garvey PB, et al. Oncologic safety and surveillance of autologous fat grafting following breast-conserving therapy. *Plast Reconstr Surg.* 2020;146:215–225.
18. Agha RA, Fowler AJ, Herlin C, Goodacre TE, Orgill DP. Use of autologous fat grafting for breast reconstruction: A systematic review with meta-analysis of oncological outcomes. *J Plast Reconstr Aesthet Surg.* 2015;68:143–161.
19. Gupta R, Brace M, Taylor SM, Bezuhly M, Hong P. In search of the optimal processing technique for fat grafting. *J Craniofac Surg.* 2015;26:94–99.
20. Groen JW, Negenborn VL, Twisk DJWR, et al. Autologous fat grafting in onco-plastic breast reconstruction: A systematic review on oncological and radiological safety, complications, volume retention and patient/surgeon satisfaction. *J Plast Reconstr Aesthet Surg.* 2016;69:742–764.
21. Kaoutzani C, Xin M, Ballard TN, et al. Autologous fat grafting after breast reconstruction in postmastectomy patients: Complications, biopsy rates, and locoregional cancer recurrence rates. *Ann Plast Surg.* 2016;76:270–275.
22. Mann RA, Ballard TNS, Brown DL, Momoh AO, Wilkins EG, Kozlow JH. Autologous fat grafting to lumpectomy defects: Complications, imaging, and biopsy rates. *J Surg Res.* 2018;231:316–322.
23. Knackstedt RW, Gatherwright J, Ataya D, Duraes EFR, Schwarz GS. Fat grafting and the palpable breast mass in implant-based breast reconstruction: Incidence and implications. *Plast Reconstr Surg.* 2019;144:265–275.
24. Rubin JP, Coon D, Zuley M, et al. Mammographic changes after fat transfer to the breast compared with changes after breast reduction: A blinded study. *Plast Reconstr Surg.* 2012;129:1029–1038.
25. Veber M, Tourasse C, Toussoun G, Moutran M, Mojallal A, Delay E. Radiographic findings after breast augmentation by autologous fat transfer. *Plast Reconstr Surg.* 2011;127:1289–1299.
26. Petit JY, Lohsiriwat V, Clough KB, et al. The oncologic outcome and immediate surgical complications of lipofilling in breast cancer patients: A multicenter study: Milan-Paris-Lyon experience of 646 lipofilling procedures. *Plast Reconstr Surg.* 2011;128:341–346.
27. Rigotti G, Marchi A, Stringhini P, et al. Determining the oncological risk of autologous lipoaspirate grafting for post-mastectomy breast reconstruction. *Aesthetic Plast Surg.* 2010;34:475–480.
28. Cohen O, Lam G, Karp N, Choi M. Determining the oncologic safety of autologous fat grafting as a reconstructive modality: An institutional review of breast cancer recurrence rates and surgical outcomes. *Plast Reconstr Surg.* 2017;140:382e–392e.
29. Xue EY, Narvaez L, Chu CK, Hanson SE. Fat processing techniques. *Semin Plast Surg.* 2020;34:11–16.
30. Ruan QZ, Rinkinen JR, Doval AF, et al. Safety profiles of fat processing techniques in autologous fat transfer for breast reconstruction. *Plast Reconstr Surg.* 2019;143:985–991.
31. Chung MT, Paik KJ, Atashroo DA, et al. Studies in fat grafting: Part I: Effects of injection technique on *in vitro* fat viability and *in vivo* volume retention. *Plast Reconstr Surg.* 2014;134:29–38.
32. Atashroo D, Raphael J, Chung MT, et al. Studies in fat grafting: Part II: Effects of injection mechanics on material properties of fat. *Plast Reconstr Surg.* 2014;134:39–46.
33. Garza RM, Paik KJ, Chung MT, et al. Studies in fat grafting: Part III: Fat grafting irradiated tissue: Improved skin quality and decreased fat graft retention. *Plast Reconstr Surg.* 2014;134:249–257.
34. Longaker MT, Aston SJ, Baker DC, Rohrich RJ. Fat transfer in 2014: What we do not know. *Plast Reconstr Surg.* 2014;133:1305–1307.