Breast augmentation: correlation between surgical planning and complication rates after surgery

Mamoplastia de aumento: correlação entre o planejamento cirúrgico e as taxas de complicações pós-operatórias

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ABSTRACT

Introduction: Breast implants remain a very popular option both for aesthetic and reconstructive plastic surgeries. A number of factors can affect the results of breast implant surgeries. The adequate planning on incision placement, need to associate mastopexy, insertion plane, and implant model increase the likelihood of adequate outcomes and reduce the need for secondary surgical treatment. This study describes the experience of a Plastic Surgery Service at the Hospital de Clínicas de Porto Alegre from 2011 to 2016 by correlating surgical planning with complication rates and surgical reintervention. Methods: A retrospective cohort that analyzed patients who underwent breast implants at the Hospital de Clínicas de Porto Alegre between 2011 and 2016, and included only cases of aesthetic breast augmentation, associated or not with mastopexy. All patients had their records analyzed, and after that an interview by phone was conducted to complement the information of the questionnaire. The analyzed was concluded with an appointment with a plastic surgeon of the service who examined the patient and confirmed the data collected for the questionnaire. No patients was evaluated by the assistant surgeon in last examination. Results: There was no significant difference between variation in surgical planning and incidence of complications. Conclusion: Still, no consensus exists regarding the best access route and plan for the breast implant. Further studies are necessary to compare the different routines of each service. Currently, best results are still based on routine systematization, precise surgical dissection and minimal contamination.

Keywords: Mamoplasty; Postoperative complications; Breast implantation.

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INTRODUCTION

Breast implants are still a popular choice for both patients of aesthetic and reconstructive plastic surgery. Breast augmentation had a 64% increase in the number of procedures performed since 2000, and it became one of the most performed aesthetic surgery in the world.

In Brazil, this is the second most common aesthetic surgery with 185,042 augmentation mammoplasties carried out in 2014\(^1\). Several studies have described both complication rates as the reoperation rates this population\(^2-4\). There are multiple factors that can affect the results of surgery with breast implant including: selection of patients, effective guiding the patient, planning for implant localization, incision location, choose of implant, surgical technique and postoperative care.

The inefficiency in any step in the treatment may result in an increased rate of complications and surgical reoperations\(^5\). Proper analysis of the physical characteristics of the patient, chest size and possible asymmetries, with the participation of the patient in the choice of the prosthesis are the most important factors in selecting the implant volume\(^6\).

The knowledge about the position of the implant and aseptic handling concepts contribute to good results and minimize the need for surgical secondary treatment\(^7\).

Patient’s satisfaction is high with this procedure, despite significant reoperation rates to treat capsular contracture, asymmetry, scar revision or other complications\(^8\).

As for surgical technique options to perform mammoplasty augmentation, there are three main decisions: the access route, anatomical plan, and type of implant used. The most popular access routes are the inframammary incision, periareolar, axillary and mastopexy associated with the inclusion of prosthesis. Regarding the plan used, the options currently accepted are subglandular, subfascial, submuscular and dual plane. Regarding the type of prosthesis, there are numerous variations. The two shapes available are anatomical and round. However, no consensus exists on what standard approach to adopt in mammoplasties augmentation, and they differ from the conducts described in the literature\(^4,8-11\).
OBJECTIVE

This study correlated surgical planning with complication rates and surgical reintervention of breast augmentation.

METHODS

This retrospective cohort study analyzed patients who underwent breast implantation at the Hospital de Clínicas/Porto Alegre between 2011 and 2016. We included only cases of aesthetic breast augmentation associated or not with mastopexy. All patients had their records reviewed, data confirmed by phone and recorded on the evaluation form (Annexes 1 and 2). The analysis was concluded with an appointment with a plastic surgeon of the service to define score of Baker and Stony Brook Scar Evaluation Scale. No patients was evaluated by the assistant surgeon in last examination.

The Stony Brook Scar Evaluation Scale was used for objective assessment of the quality of late scar. Exclusion criteria were: breast tuberous, congenital deformity of the thorax, reconstructive surgery, revision surgery or non-attendance at the evaluation of the surgeon to update the data and define the scores.

All data were typed in a Microsoft Excel 2013 spreadsheet file. We considered that all isolated augmentation mammoplasties were performed with inframammary incision or infra-areolar approach, and all implant-associated mastopexy were performed with inverted T incision. The access route techniques were correlated, as well as the dissection plans performed and the shape of the prosthesis chosen, comparing the surgical planning with the outcomes of complications and reinterventions. To analyze better the rates of major complications, reintervention and the Stony Brook Scar Evaluation Scale, data were classified by breast; total of 100 breasts.

For statistical analysis, we used the Statistical Package for Social Studies (SPSS) (IBM Corp. Released 2012; SPSS Statistics for Windows, Version 21.0 Armonk, NY. IBM Corp.). As major postoperative complications, we analyzed capsular contracture by Baker’s scale, rupture, rotation, infection, extrusion, hematoma, seroma, pain, malposition and asymmetry. The characteristics of the scar were evaluated according to the Stony Brook Scar Evaluation Scale.

Preoperative Evaluation

All patients perform preoperative exams; blood count, chest X-rays, thrombin time, activated partial thromboplastin time, electrocardiogram. Women younger than 40 years underwent breast ultrasound or mammogram if any risk factor existed or if it was not done during patient last screening tests. Other exams were requested depend on comorbidities presented by the patient.

The volume of the prosthesis is chosen according to the measures of patient’s base of breast and then discussed to the same size possibilities within limits that respect the mammary platform. The anatomical plane is chosen as the “pinch test” of patient if less than 2 cm, a double plan is chosen, and if larger than 2 cm a subglandular or subfascial is chosen. The access routes and implant shape are discussed with the patients and decided with them.

Surgical technique - access route

Before surgical marking, an inspection was carried out in the breast, anterior chest and spine in order to identify asymmetries and different heights of breast folds. The sternal midline was evident and a possible detachment area. During anesthetic induction, the antibiotic prophylaxis was done using 2 grams of cefazolin intravenously.

An incision in the mammary fold of 4 cm long with detachment 1 cm below was done to proper accommodate the prosthesis in order to avoid unsightly scars with the use of swimsuits. After detachment as planned in preoperative, the cavity was irrigated with 150 ml of saline soluction. The surgeon exchanges the gloves before proper placing the implant. Subsequently, a three-plane suture is carried out subcutaneous and subdermal with Monocryl 3.0; and intradermal with Monocryl 4.0.

In all cases we used molds to define the volume to be used. Drains were not used. Dressings used were made of micropore and sterile gauze. The patients were under observation with fixed medications, opioids, nonsteroidal anti-inflammatory drugs.

Patients were discharged 12 hours after the procedure, and instructed to take a shower only 48 hours after the surgery. Patients were followed-up every week, the dressing with micropore was maintained to define the preoperative marking for 7 days. The patient was instructed to wear a compression bra for 2 weeks.

Mastopexy associated with breast implant: Marking using A, B and C points. The Schwartzmann maneuver and de-epithelialization were done. A thin lower flap pedicle for prosthesis protection was created. In cases where there is manipulation of the mammary gland, the prosthesis was washed with solution as recommended by Adams et al., excepted the use of Bacitracin, which is not be available in the country. The de-epithelialization was carried out before test the molds (Figure 1).
Breast augmentation

Surgical technique - breast implant planning

Breast implant planning was done based on description of the literature.

RESULTS

A retrospective cohort study was carried out based on intention-to-treat approach of 50 women who underwent breast augmentation or mastopexy associated with the implant between 2011 and 2016. Patients were selected based on inclusion and exclusion criteria of previously described treatment. For better analysis, data were classified by breast; a total of 100 breasts. Patients’ age ranged from 17 years to 56 years, mean age was 36 years, a standard deviation of 10 years and a median of 36 years.

Only one patient (2%) become pregnant after plastic surgery, but no significant repercussions occurred in results. Of 6% of smokers in our sample, 33% “underwent a surgical reintervention. No rupture or rotation of the breast implant was observed. The mean volume of implants was 270 ml. Isolated mammoplasty was performed in 55% of cases (Figures 2 and 3) and mastopexy associated with implantation in 45% of cases (Figure 4).

Patients’ mean age who underwent breast augmentation was approximately 35 years, while the mean age of patients who choose to perform mastopexy associated with the implant was 39 years, with a standard deviation of 10 years in both sections.

In all cases of breast augmentation associated with mastopexy the T inverted technique was used. Patients who performed isolated breast implant, in 85% the access was via inframammary, while 15% the access was via inferior areolar. Of nine patients who underwent surgical reintervention, 5 underwent bilateral surgical reintervention (55%) and only four patients underwent unilateral surgery (45%).

When data were stratified by surgery (Figure 5), we observed that 57% of reinterventions were performed in the mastopexy associated with a prosthesis implant. In the breast augmentation group, 11% had reintervention, compared with 18% of the breasts in the mastopexy group associated with implant. Difference was not statistically significant by chi-squared test ($p = 0.487$).

Surgical reintervention was carried out in 14% of cases. Causes of reintervention were hematoma in 1% of cases, seroma 4%, asymmetry 4%, bad positioning of the prosthesis in 2% and capsular contracture in 3% breasts. The most prevalent complication was performed in 14% of patients from the implant group and in 9% of patients who underwent mastopexy associated with breast implant.

The association of dissection plan and surgical technique showed that breast augmentation by itself were similar to double plan indications (45%) and subglandular (44%). In the mastopexy associated with implant, the subglandular dissection plane was predominant (71%). Concerning correlation between dissection plane and reintervention rate (Figure 6), the subglandular plane was chosen in 56%, and required surgical reintervention in 14% of cases.
Double plan technique was performed in 32% of cases, and 19% of cases were reoperated. Subfascial plane was performed in 12% of cases and no of them required surgical reintervention. The chi-square test did not show statistical significance in the association between dissection plane and complications.

The analysis of associations between breast implant shape and reintervention rates (Figure 7) indicated the use of rounding prosthesis in 86%, which required surgical reintervention in 16% of cases. In cases with anatomical implants (14%), none required surgical reintervention. There was no statistical significance in the association between breast implant and reinterventions using the chi-square test.

Correlation between the procedure and the late aesthetic quality of the scar was assessed using a Stony Brook Scar Evaluation Scale, as mentioned above. In the breast group, 14% had grade 2, 25% grade 3, 32% grade 4 and 29% grade 5. In breast group associated with breast implant no patient had grade 1, 16% had grade 2, 30% grade 3, 13% grade 4 and 40% grade 5 (Figure 8).

No statistical significance was found in the analysis of the Stony Brook Scar Evaluation Scale in association with procedures, diagnosis, access route, dissection plane, composition, shape and texture of the prosthesis using the chi-square test. No statistical significance was found in the analysis of the Stony Brook Scar Evaluation Scale associated with age, body mass index (BMI), size from breast implant and time (in days) to return to daily life activities and practical life activities by the Kolmogorov-Smirnov and Shapiro-Wilk tests.

**DISCUSSION**

The study analyzed the effect of the surgical plan on postoperative complications rates and need of surgical reintervention in breast augmentation. It is important to perform the appropriate surgical technique to reduce complications and reoperations in mammoplasty. Complications such as capsular contracture, improper positioning and extrusion are directly related with the technical precision in which the primary procedure is done.

This study was designed to analyze specifically the effect of the surgical plan on postoperative outcome. Based on the experience of the Plastic Surgery Service of the Hospital de Clínicas de Porto Alegre. When implant
placement plans were evaluated along with reoperation rates, there were different positions in the literature. Tebbetts\(^{13}\) establishes a pattern of breast augmentation by reporting a 0% reoperation rate in 50 consecutive patients followed by 3 years.

However, other authors did not achieve the same results. Stutman et al.\(^{4}\) obtained a reoperation rate of 18% in the subglandular plan and 14% in double plan. Pereira & Sterodimas\(^{16}\) reported 2% in the subglandular plan, 6% in the double plan and 0% in subfascial. Brown\(^{17}\) described 7.2% in the subglandular and 4% in the subfascial. Montandon\(^{18}\) analyzed 546 patients and described a greater risk of general complications in submuscular (16%) than in subglandular (8%).

In this series of patients in cases that performed subglandular plan, the surgical reintervention rate was 14% and in cases of double plan this rate was 19%. In terms of prosthesis shape used, Tavares-Filho\(^{19}\) proposed an algorithm to define the choice of round or anatomical implant, this algorithm describes a higher reoperation rate in anatomical 6.2% than in the round 3.7% implants. However, in this study, the surgical reintervention rate was 16% in the round implants and no case was needed when the anatomical implants were chosen.

Regarding the comparison of access routes and postoperative outcomes, there is also no consensus in the literature. Stutman et al.\(^{4}\) did not found significant difference between the access routes and the outcomes in sample including 612 patients, however, the mammary sulcus presented a greater risk for extrusion, and the surgical reoperation rate was 24% for mammary sulcus and 13% in the periareolar route.

Authors who describe a single access route in their routine reported a lower reoperation rate. Wiener\(^{20}\) reported only 3% of reoperation cases by the mammary sulcus. Han et al.\(^{21}\), however, described 6% of reoperations in patients who underwent mammoplasty augmentation via the infra-areolar approach. Codner et al.\(^{2}\) in a retrospective analysis of 15 years on mastopexy associated with silicone implant founded a reoperation rate of 26%.

Almeida et al.\(^{21}\) in the description of a safety technique to perform mastopexy with implant reported 12% of major complications. Montandon\(^{18}\) reported a higher risk of major complications in breast augmentation associated with mastopexy (16%) than in the isolated implant (7%).

In this series, 11% of isolated breast augmentation cases were surgical reintervention, against 18% of the breasts implant-associated mastopexy group. In the evaluation of results we obtained rates similar to the large series, mainly in the studies of large centers with residency training.

As for the subfascial plane and anatomical shape implants because our sample is small, the validity of the results can be questioned. We obtained less satisfactory results regarding the late outcome of postoperative scar quality, being only 61% grade 4 or 5 in the Stony Brook Scar Evaluation Scale in the isolated breast augmentation and 53% in the breast augmentation associated with mastopexy, which motivated us to review the routine of the service.

There is still no consensus regarding the best access route and plan of breast implant, requiring studies to compare different routines of each service. Currently, the best results are still based on a systematized routine, precise surgical dissection and minimal contamination.

**CONCLUSION**

We conclude that no significant difference exist between the variation in surgical planning and the incidence of postoperative complications or need of reintervention in surgeries carried out between 2011 and 2016 in Hospital das Clínicas de Porto Alegre.

**COLLABORATIONS**

| JM | Analysis and/or interpretation of data; statistical analyses; conception and design of the study; completion of surgeries and/or experiments; writing the manuscript or critical review of its contents. |
| ACPO | Analysis and/or interpretation of data; final approval of the manuscript; writing the manuscript or critical review of its contents. |
| EL | Statistical analyses; completion of surgeries and/or experiments. |
| JB | Completion of surgeries and/or experiments. |
| CPP | Analysis and/or interpretation of data; Writing the manuscript or critical review of its contents. |
| DD | Completion of surgeries and/or experiments. |
| JH | Completion of surgeries and/or experiments. |
| MVMC | Final approval of the manuscript; conception and design of the study; writing the manuscript or critical review of its contents. |

**REFERENCES**


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**Annex 1. Data of surgical descriptions.**

<table>
<thead>
<tr>
<th>Patient's name:</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Age on the day of the surgery:</td>
<td>__________</td>
</tr>
<tr>
<td>Procedure:</td>
<td>Breast augmentation ( ) Mastopexy + Implant ( )</td>
</tr>
<tr>
<td>Diagnostic:</td>
<td>Hypoplasia ( ) Breast asymmetry ( ) Ptosis after delivery ( ) Other ptosis ( )</td>
</tr>
<tr>
<td>Pocket:</td>
<td>Submuscular ( ) Subfascial ( ) Subglandular ( ) Double plan ( )</td>
</tr>
<tr>
<td>Incision:</td>
<td>Inframammary ( ) Infra-areolar ( ) Axillary ( ) Mastopexy ( )</td>
</tr>
<tr>
<td>Prosthesis composition:</td>
<td>Jelly ( ) Saline Solution ( )</td>
</tr>
<tr>
<td>Shape of the implant:</td>
<td>Round ( ) Anatomical ( )</td>
</tr>
<tr>
<td>Projection of the implant:</td>
<td>Low ( ) Moderate ( ) High ( )</td>
</tr>
<tr>
<td>Texture of the implant:</td>
<td>Smooth surface ( ) Texture surface ( )</td>
</tr>
<tr>
<td>Size:</td>
<td>_____ ml</td>
</tr>
<tr>
<td>Time from the surgery:</td>
<td>_____ months</td>
</tr>
</tbody>
</table>

**Annex 2. Complications after surgery.**

| Contracture: | Baker grade II ( ) Baker grade III ( ) Baker grade IV ( ) |
| Rupture: | Yes ( ) No ( ) |
| Rotation: | Yes ( ) No ( ) |
| Infection: | Yes ( ) No ( ) |
| Hematoma: | Yes ( ) No ( ) |
| Seroma: | Yes ( ) No ( ) |
| Poor positioning: | Yes ( ) No ( ) |
| Assymetry: | Yes ( ) No ( ) |
| Stony Brook Scar Evaluation Scale: | 0 ( ) 1 ( ) 2 ( ) 3 ( ) 4 ( ) 5 ( ) |