Oncoplastic Breast Surgery: Is it reliable in the treatment of multifocal breast cancer? A preliminary report of a prospective randomized controlled trial

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Abstract

Background/Aim: Multifocal breast cancers (MFBCs) still have undiscoverable clinical significance. Being the standard surgical management for early breast cancer, implementation of breast-conservation therapy (BCT) as a surgical procedure for multifocal breast cancers is still questionable and needs a solid basis of clinical evidence via prospective randomized control trials.

Methods: A prospective study was conducted on female patients with operable multifocal breast cancer excluding those diagnosed with inflammatory breast cancer and those to receive neoadjuvant therapy. Surgical management was selected randomly and comprised either modified radical mastectomy (MRM) or different techniques of oncoplastic breast surgery (OPS) with a sealed envelope system based on clinical evaluation and recent guidelines for management at the Surgical Oncology Unit, Alexandria University from May 2017-May 2018. The patients were followed up until February 2021 with a median follow-up of 39 months postoperatively to assess recurrence. Analysis of different clinicopathological factors was performed to evaluate the reliability of OPS in the surgical management of MFBCs.

Results: A total of 132 patients were initially assessed for the eligibility criteria. Finally, 58 patients in the OPS group and 56 patients in the MRM group were followed up until the end of the study period. After a median follow-up of 39 months post-operatively for both groups, three patients belonging to the oncoplastic group suffered from local recurrence (5.2%). Two patients who had MRM had distant recurrence (3.6%). Although recurrence behavior was different between both groups, this was not statistically significant.

Conclusion: OPS is an oncopologically safe surgical option for selected cases of multifocal breast cancer.

Keywords: Breast cancer, Multifocal, Recurrence

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Ethics Committee Approval
Ethical committee faculty of medicine, Alexandria University, Jan 2018. The study was registered at Clinical Trial. Gov (NCT03900299). All procedures in this study involving human participants were performed in accordance with the 1964 Helsinki Declaration and its later amendments.

Conflict of Interest
No conflict of interest was declared by the authors.

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Introduction

Multifocal breast cancers (MFBCs) represent a discrete and important oncological issue. The incidence of MFBCs varies from 6% to 60% among breast cancers worldwide [1, 2]. MFBCs are defined when there are two or more synchronous cancerous foci within the same quadrant. The size of the largest tumor focus is considered for the TNM staging system [3].

MFBCs are currently encountered in the surgical oncology field more than in previous decades mostly due to the revolution in the diagnostic modalities of breast cancer. Better guidelines for their management are needed, especially regarding the optimal loco-regional therapy and their impact on survival [4-6].

Being the standard surgical management for early breast cancer, implementation of breast-conservation therapy (BCT) as a surgical procedure for MFBCs is still questionable and needs a solid basis of clinical evidence with prospective randomized control trials [7].

Conventional contraindications to BCT include any clinical conditions which may alter local control and or cosmesis. Multifocal breast cancers may be included in this category [7-10]. However, there is now growing evidence to suggest that oncoplastic conservative surgery (OPS) can be a suitable surgical procedure with acceptable local control rates [11-16].

OPS is divided into the volume-displacement procedure which includes resection with a variable volume of breast tissue, rearrangements, and mammoplasty techniques [7, 8], and volume-replacement procedures, which entails resection with immediate reconstruction using loco-regional flaps [9, 10]. In all cases, simultaneous or delayed correction in the contralateral breast can be done to achieve better symmetry [11].

The present study is directed to analyze, in a prospective series of breast cancer patients treated at a single institution, the reliability of oncoplastic breast conservative surgery as a surgical treatment for selected cases of MFBCs in terms of oncological safety.

Materials and methods

Study design and randomization

In this non-blinded two-arms parallel design randomized clinical trial, multifocal breast cancer patients were first randomly selected by the simple random sampling method. Patients were assessed for eligibility and randomly allocated at a ratio of 1:1 to either the OPS arm or MRM arm using simple randomization by tables of random number generator. Concealed allocation was achieved by sequentially numbered opaque sealed envelopes technique and by keeping the executive of the randomization unaware of study participants’ sequence.

Eligibility

Inclusion criteria: Operable female patients with MFBC diagnosed on a clinical and/or radiological basis and proved by core needle who were admitted to the Surgical Oncology Unit, Faculty of Medicine, University of Alexandria.

Exclusion criteria: Patients to receive neoadjuvant treatment or inflammatory carcinoma were excluded because of the tendency towards worse prognosis in the tumor characteristics rather than in the surgical procedure. Also, the method of localization of multifocal disease is a matter of debate [8].

Data collection

Patients diagnosed as having MFBC signed informed written consent before being enrolled in the study. They were diagnosed as MFBC on clinical and/or mammographic findings. However, MRI was resorted to whenever the mammographic findings were not conclusive (e.g., dense breast, asymmetrical architecture distortion, and lobular carcinoma). Patients were subjected to either total mastectomy or oncoplastic conservative breast surgery according to the arm of the study. Safety margin was assured in the second group by intraoperative pathological assessment. This was performed by the inking of different margins and imprinting of cytological examination of all margins by a dedicated pathologist. Effective communication between the surgeon and the attending pathologist was assured for describing the site and size of the different foci and their relation to each other and concordance with prior imaging studies. Clear margins were defined as no ink on the tumor [8].

Axillary dissection or sentinel node biopsy was performed according to the clinical and radiological states of the axilla.

Patients who underwent total mastectomy were offered different options for immediate reconstruction, only 3 (5.35%) out of 56 patients of the mastectomy accepted immediate reconstruction in the form of autologous pedicled transverse rectus abdominis muscle flap (TRAM flap) to avoid a second surgery.

The CONSORT diagram shows the flow of participants through each stage of the randomized trial (Figure 1).

After surgical intervention, the following data were recorded:

- **Tumor characteristics**: Size, nodal status, presence of lymphovascular invasion, amount of intraductal component, tumor grade, margin status, hormone receptor, and Her2 status.
- **Rate of re-excision:** If needed, was based on permanent histopathological examination of the excised specimen in patients who underwent OPS.

- **Postoperative surgical complications** or any other procedure-related problems.

All patients were scheduled for clinical follow-up every 6 months with an overall follow-up period of 39 months.

- A mammogram (or ultrasonographic examination of the mastectomy bed) and metastatic workup survey/year. The occurrence of loco-regional recurrence and distant metastases during the follow-up period were documented.

### Outcomes

1. Loco-regional recurrence or distant metastases during the follow-up period.

- Local recurrence is defined as recurrence in the original tumor bed of OPS or the mastectomy field.

- Regional recurrence refers to metastatic disease in the ipsilateral axilla or supraclavicular lymph nodes or ipsilateral involvement of internal mammary nodes.

2. The disease-free survival of patients was estimated using the Kaplan-Meier method.

### Sample size calculation

Previous studies illustrated an overall survival of 94% vs. 90% among patients undergoing breast conservation relative to the group treated with mastectomy, respectively, with no significant difference [17]. If there is truly no difference between the standard and experimental treatment (around 94% in both groups), then 110 patients (55 per each intervention arm) are needed to be 95% sure that the limits of a two-sided 95% confidence interval will exclude a difference between the standard and experimental group of more than 15%. The calculation was based on the following formula for equivalence trial to achieve 80% power at 0.05 significance level:

\[
 n = \frac{2 \times f(\alpha, \beta) \times \pi \times (100 - \pi)}{d^2}
\]

where \(\pi\) is the true percent “success” in both the control and experimental treatment groups, and

\[
f(\alpha, \beta) = \Phi^{-1}(1-\alpha) + \Phi^{-1}(\beta/2)
\]

where \(\Phi^{-1}\) is the cumulative distribution function of a standardized normal deviate. We anticipated a potential loss of follow-up of 10%. To account for this, we initially enrolled 132 patients in the trial.

### Ethical consideration

The study was approved by the ethical committee of the Faculty of Medicine, University of Alexandria. Also, the study was registered in clinicaltrials.gov (NCT03900299).

https://clinicaltrials.gov/ct2/show/NCT03900299?recrs=d&cond=Breast+Cancer&cntry=EG&draw=2&rank=2

The patients’ records were kept confidential and all signed informed consent forms.

### Statistical analysis

Data are expressed as mean (SD), and numbers or percentages where appropriate. The Chi-square test was performed to study the significant associations between categorical variables. Fischer exact significance, as well as Monte-Carlo significance, were used if more than 20% of the total expected cell counts <5 at 0.05 level of significance. Kaplan-Meier survival analysis was conducted to compare overall survival and recurrence-free survival outcomes between the two interventions using the Log-rank test. All statistical tests were judged at a 0.05 significance level using IBM SPSS statistics program version 21.

### Results

A total of 132 patients were initially assessed for eligibility. Twelve patients were excluded, five refused to participate after enrollment, and 7 patients did not meet the eligibility criteria. Sixty patients were randomized, and 2 patients in the OPS group and 4 patients in the MRM group were lost to follow-up. Finally, 58 patients who were subjected to OPS and 56 patients who underwent MRM continued follow-up (CONSORT diagram).

A summary of the clinicopathological characteristics of patients is given in Table 1. Both groups were similar regarding their age, stage, tumor size, histopathological type, grade, and lymphovascular invasion, presence of excessive intraductal component, hormonal status & Her2 expression.

The adjuvant treatments administered in both groups, whether chemotherapy, radiotherapy, or hormonal, is illustrated in Table 1.

### Table 1: A summary of the clinicopathological characteristics of patients

| Item                  | OPS (n=58) | MRM (n=56) | \( \chi^2 \) | P-value |
|-----------------------|------------|------------|---------------|---------|
| Age in years          |            |            |               |         |
| <40                   | 18         | 10         | 4.011         | 0.026   |
| 40-                  | 18         | 10         |               |         |
| 50-                  | 14         | 14         |               |         |
| 50+                  | 21         | 29         |               |         |
| T staging             |            |            |               |         |
| T1                   | 24         | 31         | 4.139         | 0.007   |
| T2                   | 21         | 22         |               |         |
| T3                   | 0          | 1          |               |         |
| N staging             |            |            |               |         |
| N0                   | 32         | 26         | 0.872         | 0.351   |
| N1                   | 26         | 44         |               |         |
| N2                   | 0          | 0          |               |         |
| STAGE                 |            |            |               |         |
| I                    | 21         | 32         | 3.773         | 0.100   |
| II                   | 37         | 63         |               |         |
| III                  | 0          | 1          |               |         |
| Histopathology        |            |            |               |         |
| IDC                  | 58         | 100        | 1.045         | 0.491   |
| Mucoid               | 0          | 0          |               |         |
| GRADE                | III        | 57         | 100.0        | 0.947   |
| II                   | 57         | 98.3       |               | 1.000   |
| I                    | 1          | 1.7        |               |         |
| Estrogen              | -ve        | 6          | 5.794         | 0.123   |
| +                    | 5          | 8.6       |               |         |
| ++                   | 18         | 31.0      |               |         |
| +++                  | 29         | 50.0      |               |         |
| Progesterone          | -ve        | 7          | 12.5         | 0.054   |
| +                    | 10         | 17.2      |               |         |
| ++                   | 32         | 55.2      |               |         |
| +++                  | 9          | 15.5      |               |         |
| HER2                 | -ve        | 54         | 93.1         | 0.555   |
| +                    | 3          | 1.7       |               |         |
| ++                   | 1          | 1.7       |               |         |
| +++++                | 2          | 3.4       |               |         |
| Intra ductal component| No         | 47         | 81.0         | 0.239   |
| Yes                  | 11         | 19.0      |               | 0.070   |
| LV1                  | No         | 26         | 44.8         | 0.306   |
| Yes                  | 48         | 52.0      |               | 0.580   |
| POS CHM              | No         | 32         | 55.2         | 2.108   |
| Yes                  | 20         | 36.2      |               | 0.001   |
| POS XRT              | No         | 58         | 100.0        | 5.794   |
| Yes                  | 10         | 19.6      |               | 0.001   |
| POS HOR              | No         | 58         | 100.0        | 0.307   |
| Yes                  | 2         | 1.7       |               | 1.000   |

\( \chi^2 \) = Chi-square test, \( \text{Fisher exact probability} \), \( P<0.01 \) (highly significant), \( MC \) = Monte Carlo exact probability, \( ** P<0.05 \) (significant)

### Regarding patients who had OPS: Various oncoplastic techniques were utilized (Table 2). Lateral therapeutic mammoplasty was the commonest technique adopted; it was used in 20 patients (34.48%) who had tumors located in the upper outer quadrant (Figure 2).
Table 2: Oncoplastic techniques used in the study

| Technique                  | Tumor location          | Number of cases |
|----------------------------|-------------------------|-----------------|
| Lateral therapeutic mammoplasty | Upper outer quadrant | 20(34.48%) |
| Round block                | Upper outer quadrant    | 14(24.13%)     |
|                           | Lower inner quadrant    | 5(8.62%)       |
|                           | Upper inner quadrant    | 5(8.62%)       |
|                           | Upper pole              | 1(1.72%)       |
| Parallelogram lumpectomy   | Upper outer quadrant    | 9(15.51%)      |
|                           | Upper inner quadrant    | 3(5.17%)       |
| J therapeutic mammoplasty  | Lower pole              | 4(6.89%)       |
| V therapeutic mammoplasty  | Lower outer quadrant    | 4(6.89%)       |
| LD FLAP                    | Upper outer quadrant    | 3(5.17%)       |
| Reduction therapeutic mammoplasty | Upper outer quadrant | 2(3.44%)     |
| Grisetti flap              | Central (retro areolar) | 2(3.44%)       |

Figure 2: Pre-operative markings of therapeutic lateral mammoplasty and outcome after the remodeling of breast tissue.

Five patients (8.6%) required re-excision after frozen section examination of the excised specimen margins. The results of the margin assessment were confirmed on the final examination of the permanent paraffin block.

Postoperative complications were encountered in 33.92% and 31.03% in the MRM & OPS groups, respectively. They were diverse, related to the nature of the procedure performed, and statistically not significant with \(P=0.90\) for early complications and \(P=0.57\) for late complications (Table 3).

Table 3: Postoperative complications after OPS and MRM

| Complications          | MRM (n=56) | OPS (n=58) | \(P\)-value |
|------------------------|------------|------------|-------------|
| Early complications    |            |            |             |
| Hematoma               | 2(3.57%)   | 1(1.72%)   | 0.900       |
| Seroma                 | 11(19.64%) | 8(13.79%)  |             |
| Abscess                | 1(1.78%)   | 2(3.44%)   |             |
| Skin or Flap necrosis  | 2(3.57%)   | 2(3.44%)   |             |
| Total no of complications | 16(28.57%) | 13(22.41%) |             |
| No complications       | 40(71.42%) | 45(77.59%) |             |
| Late complications     |            |            |             |
| Scar fibrosis          | 1(1.78%)   | 1(1.72%)   |             |
| Keloid                 | 2(3.57%)   | 2(3.44%)   |             |
| Steatonecrosis         | 0          | 2(3.44%)   |             |
| Total no of complications | 8(5.35%)  | 5(8.62%)   |             |
| No complications       | 53(94.65%) | 53(91.38%) |             |

After a median follow-up period of 39 months, three patients belonging to the oncoplastic group suffered from local recurrence (5.2%) and two patients who had MRM had distant recurrence (3.6%). Although the recurrence pattern differed between the two groups, local recurrence, distant recurrence, and the overall recurrence rates were not significant (\(P=0.244\), \(P=0.239\), \(P=0.671\), HR: 1.474 at 95% CI, respectively) (Table 4, 5).

### Table 4: univariate analysis of local and distant recurrence in both groups

| Item          | Surgery | MRM | \(X^2\) | \(P\)-value |
|--------------|---------|-----|---------|-------------|
| Recurrence    | No      | 54  | 96.4%   |             |
|              | Yes     | 5   | 94.8%   |             |
| Local         | No      | 54  | 96.4%   |             |
|              | Yes     | 5   | 94.8%   |             |
| Distant       | No      | 2   | 3.6%    |             |
|              | Yes     | 13  | 22.41%  |             |

### Table 5: Rate of recurrence in both groups

| No Recurrence | Recurrence | \(P\)-value | HR (95%C.I) |
|---------------|------------|-------------|-------------|
| No            | 55         | 94.8%       |             |
| Yes           | 3          | 5.2%        | 0.671       |

The Kaplan-Meier method revealed the disease-free survival (DFS) to be 94.8% and 96.4% in OPS & MRM groups, respectively. This difference was not significant (Figure 3, Table 6).

### Table 6: Kaplan-Meier survival analysis for disease-free survival with surgery

| Surgery | Mean | % 1 year | % End of study | Log-rank \(X^2\) | \(P\)-value |
|---------|------|----------|----------------|----------------|-------------|
| OPS     | 57.57| 96.6     | 94.8           |                 |             |
| MRM     | 58.50| 96.4     |                | 0.183           | 0.668       |

Figure 3: Kaplan-Meier survival curve for disease-free survival with surgery

Discussion

Multifocal breast cancer is an entity that needs consideration. Its biological basis is not well understood, especially whether it is the result of a simultaneous overgrowth of tumor foci or an outcome of extensive intraductal carcinoma [19]. Many studies found it to have a different clinical outcome with a poorer prognosis if compared with unifocal breast cancer of the same stage [20] with This is attributed to the high invasive tumor burden and association with lymph node metastases [21]. The reported association between multifocality and the outcome of patients in early breast cancer is variable [20, 22, 23]. It has a higher possibility of recurrence and adverse survival after surgical treatment, especially if breast conservative surgery is performed. This paradox directed many surgeons to perform mastectomy for these patients [24, 25]. This was the reported trend in the past two decades [9, 10, 24, 26]. Recently, some studies found a lower incidence of local recurrence in patients with multifocal breast cancer treated with breast conservative surgery [27, 28]. However, most were retrospective.

Thus, the present study was conducted prospectively to compare the results of mastectomy and breast conservative therapy in the treatment of multifocal breast cancer. We included only MFBC patients diagnosed on a clinical and radiological basis. The pattern of recurrence was different between both groups; local recurrence only occurred in patients who...
underwent OPS while patients who underwent MRM suffered only from distant recurrence. The overall recurrence rate and the disease-free survival (DFS) were not significantly different between the two groups.

Reviewing the literature, few studies were found investigating breast conservative therapy (BCT) and mastectomy as treatment options for MFBC. The majority found no difference between both modalities regarding local recurrence (LR) and disease-free survival (DFS) [17, 29]. Moreover, Wolters et al. found that BCT and mastectomy are suitable comparable choices in the surgical treatment of T1/2 MFBC [14]. These previous results are in agreement with our work. One of the strengths of our study is that we determined the eligibility criteria for inclusion and exclusion of patients like everyday clinical practice while applying both interventions, so the findings from this study could be generalizable. We encountered the most clinically important outcomes, and we also considered the potential complications after surgery.

Although the follow-up period in our study is relatively short, the paucity of prospective RCT addressing MFBC encouraged the authors to publish the results as a preliminary study with early results. Still, longer follow-up periods are needed. However, this study has the advantage of being a prospective one.

Conclusions

From the present work, we can conclude that breast OPS is a safe option to treat selected cases of MFBC that can, nowadays, is better characterized with the wide use of magnetic resonance mammography. It achieves a wide resection margin enabling the removal of all tumor foci.

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