Repeat Breast-Conserving Surgery Versus Salvage Mastectomy for Ipsilateral Breast Tumour Recurrence After Breast-Conserving Surgery in Breast Cancer Patients: A Meta-Analysis

Caiqin Mo†, Weihong Ruan†, Junyu Lin†, Huaying Chen† and Xiangjin Chen†*

†Department of Breast and Thyroid Surgery, The First Affiliated Hospital of Fujian Medical University, Fujian, China,
†Department of General Surgery, Traditional Chinese Hospital of Xiamen, Fujian, China, ‡The First Clinical Medical College, Fujian Medical University, Fujian, China

**Correspondence:** Xiangjin Chen
rjbhcxj@fjmu.edu.cn
orcid.org/0000-0002-0132-6673

†These authors share first authorship

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**Background:** Salvage mastectomy (SM) is the standard surgery for ipsilateral breast tumour recurrence (IBTR). However, whether repeat breast-conserving surgery (RBCS) is an alternative method remains unclear. We performed a meta-analysis to compare the effects of RBCS and SM after IBTR for breast-conserving surgery (BCS).

**Methods:** We searched PubMed, Cochrane, Wiley Online and Embase for controlled studies comparing RBCS and SM after IBTR for BCS (published between 1993 and 2019, published in English). Our main endpoints were the secondary local recurrence rate (SLRR), distant metastasis rate (DMR) and overall survival (OS). We used a random-effects model or fixed-effects model for data pooling.

**Results:** Fifteen of the 424 eligible studies were ultimately included, and all studies were retrospective cohort studies (n=2532 participants). 1) SLRR: The SLRR of RBCS was higher than SM (pooled relative rate (pRR) = 1.87, 95% CI 1.22 - 2.86, P=0.004). Stratified analysis was performed according to whether radiotherapy was performed after salvage surgery (radiotherapy group: 2ndRT, no radiotherapy group: no-2ndRT), and the following results were revealed: pRR=0.43 (95% CI 0.20-0.95, P=0.04) for group 2ndRT; and pRR=2.30 (95% CI 1.72-3.06, P<0.00001) for group no-2ndRT. These results showed that the main cause of heterogeneity was salvage radiotherapy. 2) DMR: No significant difference in the DMR was observed between RBCS and SM (pRR = 0.61, 95% CI 0.37 - 1.01, P=0.05). 3) OS: No significant difference in OS was observed between RBCS and SM (pRR=0.65, 95% CI 0.39 - 1.08, P=0.10).
INTRODUCTION

Breast-conserving surgery (BCS) is a standard surgical method for early breast cancer. However, local recurrence exists. Ipsilateral breast tumor recurrence (IBTR) is defined as the reappearance of breast cancer in the region of the ipsilateral breast/chest wall or the draining regional lymph node basins (1). The 10-year ITBR rate is approximately 5-10% (2). For IBTR, approximately 6-7% of all patients have inoperable disease (3, 4), and 5-10% develop distant metastasis simultaneously (5, 6).

Opportunities for the detection of small and isolated IBTR diagnoses have increased (7), and the demands of repeat breast-conserving surgery (RBCS) have become more urgent. However, salvage mastectomy (SM) is the standard surgical method for operable IBTR (2, 8, 9), and whether RBCS is an alternative method in patients with IBTR is controversial. Some studies reported that the prognosis of RBCS was worse than SM (4, 10–12), but other studies have not (13–19). Therefore, we performed a meta-analysis of the secondary local recurrence rate (SLRR), distant metastasis rate (DMR) and overall survival (OS) of RBCS or SM in IBTR patients after BCS to further evaluate the feasibility of RBCS for IBTR after BCS.

MATERIALS AND METHODS

Search Strategy

This meta-analysis is reported according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA). We selected 15 relevant studies published between 1988 and 2019 after searching Embase, PubMed, Cochrane, and Wiley Online databases (only published in English). We also searched the reference lists of important articles manually. The complete search strategy of PubMed is shown in Data S1.

Selection and Data Extraction

The inclusion criteria were as follows: (a) studies comparing RBCS and SM of IBTR after BCS, regardless of whether radiotherapy was administered after the first BCS; (b) retrospective cohort studies or prospective cohort studies; and (c) studies that included data on the SLRR, DMR or OS.

The exclusion criteria were as follows: (a) unreasonable research design, incomplete data, or unclear endpoints; (b) poor data sources or sources from the same center; (c) data from the SEER database; and (d) studies not published in English.

Quality Assessment

Two independent investigators reviewed the study titles and abstracts independently and extracted and analyzed the data, and disagreements were resolved by a third investigator. We extracted the following data: sample size, inclusion time, follow-up time, number/rate of secondary recurrence events, number/rate of distant metastatic events, number/rate of time of death, radiotherapy information and other treatment information. The risk for bias according to the PRISMA recommendations were assessed by two independent reviewers.

The Newcastle-Ottawa Scale (NOS) was used to assess the quality of the included literature. A higher score indicated a lower risk of bias. An overall NOS score ≥ 6 was considered acceptable (Data S1).

Statistics Analyses

The following outcomes were assessed: SLRR, DMR and OS. We analyzed the SLRR, DMR and OS as binary count variables. If the 5-year and 10-year OS were provided, the 10-year OS was used as the final data. The overall relative risk (RR) was calculated and the Cochran Q test was used to assess heterogeneity between studies. If testing was performed to assess the magnitude of the heterogeneity between studies, and if values was greater than 50%, moderate-to-high heterogeneity was indicated. A fixed-effect model was used for low heterogeneity, and a random-effect model was used for moderate-to-high heterogeneity. To confirm the effect of radiotherapy after RBCS on recurrence and survival, stratified analysis was performed for the SLRR based on whether radiotherapy was administered after RBCS.

We assessed the possibility of publication bias by funnel plots. We assessed funnel plot asymmetry by using Begg’s and Egger’s tests and defined significant publication bias as two-tailed p < 0.05. Sensitivity analysis was performed by omitting one study. We used Stata (version 12.0) and Endnote X9 for all statistical analyses.

RESULTS

Study Selection Results

A total of 424 studies were identified. No prospective randomized controlled studies were included. Fifteen studies (with 2532 participants) were ultimately included in our meta-analysis (Figure 1). The lowest NOS score of included studies is 7 and the detailed NOS study equality evaluation via the

Conclusions: The SLRR of RBCS was higher than SM for ITBR after BCS, but survival was not affected. RBCS may be used as an alternative for IBTR patients after BCS with strict control for several indications, such as tumor size, recurrence interval and biological behavior, and attaching importance to subsequent salvage radiotherapy and systematic therapy.

Keywords: meta-analysis, repeat breast-conserving surgery, salvage mastectomy, ipsilateral breast tumor recurrence, breast cancer
Newcastle-Ottawa Scale was shown in the supplementary materials Data S1.

**Study Characteristics**

All studies were retrospective cohorts published between 1988 and 2019. The median follow-up time ranged from 52 months to 20 years. Of the 2532 participants, 633 underwent RBCS, and 1899 underwent SM. Our main endpoints were the SLRR, DMR and OS. Thirteen studies reported the SLRR, 4 studies reported the DMR, and 7 studies reported OS. Systematic data from the studies by Alpert, T.E. et al., Chen, S.L. et al. and Mccready et al. were not available, and the other 12 studies received certain systematic treatment. See Table 1 for the detailed characteristics of the included studies.

**Meta-Analysis Results**

1. **SLRR**

A total of 13 studies reported the SLRR. Among these studies, 431 patients underwent RBCS, and 1224 patients underwent SM ($I^2=56\%$, $P=0.007$), which suggests heterogeneity among the studies. A random-effects model was used, and the combined effect size of $pRR=1.87 (1.22-2.86)$, $P=0.004$ (Figure 2) suggested that the SLRR of RBCS was significantly higher than SM. Stratified analysis was performed according to whether radiotherapy was performed after salvage surgery to examine sources of heterogeneity (Figure 3). The results revealed that radiotherapy after RBCS (2ndRT group) was performed in 2 studies, including 58 patients in the RBCS group and 189 patients in the SM group. Eleven studies did not perform radiotherapy after surgery (no-2ndRT), including 373 patients in the RBCS group and 1035 in the SM group. After stratification, there was no significant heterogeneity in the two groups (for 2ndRT, $I^2=0\%$, $P=0.38$, and for no-2ndRT, $I^2=16\%$, $P=0.29$). A fixed-effect model was used for stratification, and the results revealed a $pRR$ of 0.43 for 2ndRT (95% CI 0.20-0.95, $P=0.04$) and a $pRR$ of 2.30 for no-2ndRT (95% 1.72-3.06, $P < 0.00001$) (See the funnel plot in Figure 4). The results of Begg’s test ($P=0.951$) and Egger’s test ($P=0.823$) suggested no publication bias. Sensitivity analysis was performed by omitting one study (Figure 5). Removal of either study showed no significant effect on $pRR$.

2. **DMR**

Four studies (with 176 RBCS and 373 SM participants) were included in the DMR analysis. The results suggested heterogeneity among the studies ($I^2=56\%$, $P=0.08$), and we used a random-effects model. The combined effect size [$pRR=0.61 (95\% 0.37-1.01)$, $P=0.05$ (Figure 6)] showed no significant difference between RBCS and SM in DMR. The results of Begg’s test ($P=1.0$) and Egger’s test ($P=0.747$) showed no publication bias. Sensitivity analysis was performed using the...
method of omitting one study (Figure 7), and no significant changes in pRR were observed.

3. OS
Eight studies (with RBCS 337 and 867 participants) were included in the analysis of OS. The results suggested heterogeneity among the studies ($I^2=87\%$, $P<0.00001$), and we used a random-effects model. The combined effect size [pRR=0.65, 95% CI 0.39-1.08, $P=0.10$ (Figure 8)] showed no significant difference between RBCS and SM in OS. The results of Begg’s test ($P=1.0$) and Egger’s test ($P=0.069$) showed no publication bias. Sensitivity analysis was performed by omitting

$TABLE 1 | Characteristics of included studies.$

| Author                  | Published year | Country | Included time | Radiotherapy after primary breast-conserving | Radiotherapy after repeated breast-conserving | Endpoint | Systematic treatment | Median follow-up time (N) |
|-------------------------|----------------|---------|---------------|---------------------------------------------|---------------------------------------------|----------|----------------------|--------------------------|
| Abner, A.L. (20)        | 1993           | US      | 1968-1985     | Yes                                        | No                                          | SLRR     | Yes                  | 79 (5-233) months (16 123) |
| Alpert, T.E (13)        | 2005           | US      | Before 1999   | Yes                                        | No                                          | SLRR, DMR, OS | Yes           | 13 (9-19) years (30 116) |
| Chen, S.L (10)          | 2008           | US      | 1988-2004     | Yes                                        | No                                          | OS       | No                   | 10 years (179) 567       |
| Dalberg, K. (6)         | 1998           | Switzerland | 1976-1985     | Yes, 79%                                   | No                                          | SLRR     | Yes                  | 13 (9-19) years (14 65)  |
| Fodor, J (14)           | 2008           | Hungary | 1983-1987     | Yes, 50%                                   | Yes, only 4                                  | SLRR, DMR, OS | Yes           | 165 (75-240) months (32 32) |
| Komoke, Y (19)          | 2005           | Japan   | 1986-1993     | Yes                                        | No                                          | SLRR     | Yes                  | 20 years (55 88)        |
| Kurtz, J, M (21)        | 1988           | Switzerland | before 1985   | Yes                                        | No                                          | SLRR     | Yes                  | 7 (1-20) years (52 66)   |
| Kurtz, J, M (22)        | 1990           | France  | 1963-1982     | Yes                                        | No                                          | SLRR, DMR | Yes           | 11 (5-24) years (34 36) |
| Lee, J.H (23)           | 2015           | Korea   | 1955-2011     | Yes                                        | Yes, partial                                | SLRR, DMR | Yes           | NS (23 108)             |
| McCready (24)           | 1994           | Canada  | 1977-1986     | No                                         | Yes                                        | SLRR     | Yes                  | 6 years (19 33)          |
| Salvadori, B (4)        | 1999           | Italy   | 1970-1989     | Yes                                        | clear                                       | SLRR, DMR, OS | Yes           | 73 (1-192) months (57 133) |
| Sellam (25)             | 2019           | Israel  | 1987-2014     | Yes                                        | Yes, only 3                                  | SLRR, DMR | Yes           | 14 (1-30) years (47 74)  |
| Smajkovic, V (26)       | 2019           | Switzerland | 1999-2015    | Yes                                        | No                                          | SLRR     | Yes                  | 59 (1-189) months (39 156) |
| Voogd, A.C (27)         | 1999           | Netherlands | 1980-1992   | Yes                                        | No                                          | OS       | Yes                   | 56 (2-189) months (20 229) |
| Wapnr, I.L (12)         | 2017           | Switzerland | 2003-2010    | Yes                                        | Yes, only 3                                  | SLRR     | Yes                  | 4.9 years (16 73)        |

SLRR, secondary local recurrence rate; DMR, distant metastasis rate; OS, overall survival; RBCS, repeated breast-conserving surgery; SM, salvage mastectomy.

FIGURE 2 | Forest plot of repeated breast-conserving surgery (RBCS) versus salvage mastectomy (SM) after ipsilateral breast tumor recurrence (IBTR), comparing secondary local recurrence rate (SLRR). The meta-analysis was performed with random effects model. RR more than 1 means the results favor SM group.
one study (Figure 9), and no significant changes in pRR were observed.

**DISCUSSION**

Since there is still some risk of recurrence after BCS, these patients face decisions of whether to undergo RBPS. For patients who choose breast-conservation at the initial diagnosis, there must be a certain willingness to preserve the breast for cosmetic or trauma reasons. Therefore, there are always some patients who desire RBPS after IBTR. However, no prospective studies compared the therapeutic effects of RBPS and SM after IBTR. This meta-analysis was performed to determine the safety of RBPS relative to SM in terms of SLRR, DMR, and OS.

The results of previous studies on whether RBPS was better than SM after IBTR varied greatly. Some studies reported that RBPS was not worse than SM (12, 13, 15, 21, 26, 28), and some studies reported the opposite results (8, 10, 11, 14). Other studies asserted that it did not suggest a worse prognosis, although the SLRR of RBPS was higher (25), which is consistent with our conclusions. The SLRR of RBPS was approximately 7-31% (8, 13, 14, 29, 30), and the SLRR of SM was approximately 3-32% (8, 13, 21). Nevertheless, these data were based on studies that did not offer secondary radiotherapy and were heterogeneous between groups. Therefore, real-world data must be further refined.

The SLRR of RBPS was initially higher than SM after IBTR (pRR = 1.87, 95% CI 1.22-2.86, P=0.004). The stratification analysis results showed that the heterogeneity came from the delivery of secondary radiotherapy, which suggests that supplementary radiotherapy after RBPS plays an important role in improving the local control rate. However, there were only two studies that explicitly offered secondary radiotherapy in our study. Previous studies also supported the opinion that secondary radiotherapy after salvage surgery reduces the SLRR (13, 25, 26, 31-38). Su Y et al. (11) reported a worse prognosis after RBPS, poor OS (HR=1.522, 95% CI 1.317-1.759), and poor breast cancer-specific survival (BCSS) (HR =1.666, 95% CI 1.319-2.105). However, for patients with radiotherapy after RBPS, the mortality was comparable to SM regardless of whether radiotherapy was administered after the first breast-conserving surgery, although BCSS was worse for RBPS patients undergoing secondary radiotherapy (HR=1.54, 95% CI 1.037-2.286, p=0.032). Therefore, radiotherapy plays an important role in improving the prognosis after RBPS. However, not all studies agree (25).

Tolerance should also be considered for decisions on secondary radiotherapy after salvage surgery. Some studies reported that a full dose of whole-breast radiotherapy was not tolerated and led to unacceptable toxicity and cosmetic damage. In reality, 75.3% of patients do not receive this radiotherapy (39). More studies showed that supplementary radiotherapy was safe, the side effects were tolerated, and good cosmetic effects were achieved at the same time due to improvements in radiotherapy technology (40-42). Secondary radiation after RBPS may include high-dose external radiation (43), 3-dimensional conformal partial-breast reirradiation (40) and perioperative brachytherapy (26). Hannoun-Levi et al. (31) and Chadha et al. (41) used low-dose-rate multi-catheter implants at centres with considerable brachytherapy experience, and both studies reported excellent outcomes for local control and toxicity. Arthur et al. used external-beam conformal partial-

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**FIGURE 3** | Forest plot of stratification analysis of repeated breast-conserving surgery (RBPS) versus salvage mastectomy (SM) after ipsilateral breast tumor recurrence (IBTR), comparing secondary local recurrence rate (SLRR). The meta-analysis was performed with fixed-effects model. RR more than 1 means the results favor SM group. 2ndRT: radiotherapy was performed after salvage surgery; no-2ndRT: radiotherapy was not performed after salvage surgery.
breast radiotherapy (PBI) in RTOG 1014 trial. The Radiation Therapy Oncology Group (RTOG) 1014 trial was the most important prospective study about partial-breast radiotherapy (PBI) after RBCS. One year toxicity report of RTOG 1014 trial was published at a median follow-up of 3.64 years in 2017 (40), there were 4 patients (6.9%) with late grade 3 treatment-related adverse events. Data about effectiveness of treatment and adverse events was updated in 2020 after 5.5 years follow-up (44), four patients (7%) had grade 3 and none had grade 4 adverse events. The 5-year cumulative recurrence of patients who underwent RBCS plus 3D-PBI was 5% (95%CI, 1%-13%) and patients underwent ipsilateral mastectomies was 10% (95%CI, 4%-20%). Both distant metastasis-free survival and overall survival rates were 95% (95%CI, 85%-98%). A study of the GEC-ESTRO Breast Cancer Working Group showed that accelerated PBI with interstitial brachytherapy is feasible and effective in preventing second local recurrence and the OS is at least equivalent to those performed with SM (32). 217 IBTR patients who accepted BCS and whole breast radiation (WBI) were included, the patients were performed with accelerated PBI with interstitial brachytherapy after RBCS, 5 and 10-year SLRR were 5.6% (95% CI: 1.5%–9.5%) and 7.2% (95% CI: 2.1%–12.1%), 5 and 10-year DM were 9.6% (95% CI: 5.7%–15.2%) and 19.1% (95% CI: 7.8%–28.3%), and 5 and 10-year OS were 88.7% (95% CI:
83.1%–94.8%) and 76.4% (95% CI: 66.9%–87.3%). G3-4 complication rate was 11%. Therefore, PBI may be a good re-irradiation method which can provide good therapeutic effectiveness, tolerability and aesthetics after RBCS, which was supported by many other studies (36, 37). To ensure the safety of the second radiotherapy and cosmetic effects, intervention time (perioperative or intraoperative), radiotherapy range (partial or total), and precision (3D conformal) should be considered cautiously.

The results showed no significant difference in the DMR or OS between RBCS and SM after IBTR (DMR: pRR=0.61 (95% CI 0.37-1.01), P=0.05; OS: pRR=0.65 (95% CI 0.39-1.08), P=0.10). No publication bias or differences in sensitivity were observed. Therefore, no significant difference in prognosis was observed in our study despite the lower SLRR of RBCS after IBTR than SM. This conclusion is similar to Sellam Y et al. (25).

An increasing number of recent studies used RBCS (10, 14, 29, 30, 45). The selection of a suitable population for RBCS is very important. The German Society of Radiation Oncology (DEGRO) expert panel guidelines published indications for RBCS in 2016, namely, single disease, size <3 cm, age > 50 years, treatment-free interval (TFI) >48 months, and patient willingness (46). Therefore, some studies suggested that the key factor affecting the prognosis of IBTR is not the method of salvage surgery but the biological behavior of the tumor. IBTR, which has good biological behavior and may be detected early, is suitable for RBCS, but it is not suitable for patients with BRCA mutations (13). It has also been suggested that oestrogen receptor (ER)-positive status and subsequent endocrine therapy should be emphasized (23). From the tumor biological behavior perspective, studies suggested the division of ITBR into two categories: true recurrence (TR) and new primary (NP) (47). The criteria for differentiating TR and NP are location of recurrence, the positive margin of the primary tumor, and pathological characteristics (15). For NP, the DFI was longer, the patients were younger, and the tumor was generally in different quadrants compared to the primary tumor, and these patients had a better survival rate (15, 48). TFI is the most frequently reported key prognostic factor, and it reflects the biological behavior of ITBR (11, 14, 19, 23, 49, 50).

In addition to secondary radiotherapy after salvage surgery, systematic treatment is also important for prognosis. In chemotherapy for isolated locoregional recurrence of breast cancer (CALOR), a study was performed to define the
significance of systemic chemotherapy after recurrence for single and operable IBTRs, and the results indicated that the hazard ratio (HR) of the risk of recurrence after salvage surgery between the chemotherapy and no-chemotherapy group was 0.59, and the SM was not a major factor affecting survival (51). Other studies also confirmed the importance of systemic therapy (12, 23).

Our analyses have the following limitations (1). Due to the absence of RCTs, conditions were not balanced between groups, and the SM group had a greater tumor load and later tumor staging (10, 13, 14, 21). Only three studies were basically balanced at baseline (12, 15, 26). However, the SLRR of the RBCS group included in the literature was higher despite the lower tumor load. Therefore, we speculated that the SLRR of RBCS was higher than the real world (2). The time span of the included studies was too long (1993-2019) because the therapeutic effects of breast cancer, especially systemic treatment, have made great progress in recent years, which could cause some deviation.

In conclusion, the SLRR of RBCS is higher for IBTR patients after BCS, but it does not affect survival. Relatively more studies reported that the SLRR was higher after RBCS than after SM, and more studies supported SM. However, due to advancements in radiotherapy technology and systematic treatment, the recurrence rate in the real world may not necessarily be higher. Indications for RBCS must be strictly controlled, namely, tumor size, number of recurrent tumor, lymph node invasion, TFI, age, and biological behavior (such as ER expression, HER2 expression, and BRCA mutation). The importance of secondary radiotherapy and systematic treatment should be emphasized. Methods to avoid the overtreatment of low-risk patients and provide adequate treatment to high-risk patients should be the focus and direction of future research.

FIGURE 8 | Forest plot of repeated breast-conserving surgery (RBCS) versus salvage mastectomy (SM) after ipsilateral breast tumor recurrence (IBTR), comparing overall survival (OS). The meta-analyse was performed with random effects model. RR more than 1 means the results favor SM group.

FIGURE 9 | Sensitivity analysis of meta-analysis of overall survival (OS) by the method omitting one study.
DATA AVAILABILITY STATEMENT
The original contributions presented in the study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

AUTHOR CONTRIBUTIONS
CM, WR, and JL: Paper flow Design, Data collection and extraction, Write the manuscript, Analysis of the results. CM and HC: Statistics, Assess the bias and quality, XC: Provide the idea of article, Make important revisions to the paper. All the authors agree the statement above. All authors contributed to the article and approved the submitted version.

REFERENCES
1. Wapnir IL, Khan A. Current Strategies for the Management of Locoregional Breast Cancer Recurrence. Oncol (Williston Park) (2019) 33:19–25.
2. Wapnir IL, Anderson SJ, Mamounas EP, Geyer CE, Jeong JH, Tan-Chiu E, et al. Prognosis After Ipsilateral Breast Tumor Recurrence and Locoregional Recurrences in Five National Surgical Adjuvant Breast and Bowel Project Node-Positive Adjuvant Breast Cancer Trials. J Clin Oncol (2006) 24:2028–37. doi: 10.1200/JCO.2005.43.3273
3. Kurtz JM, Amalric R, Brandone H, Ayme Y, Jacquierme J, Pietra JC, et al. Local Recurrence After Breast-Conserving Surgery and Radiotherapy. Frequency, Time Course, and Prognosis. Cancer (1989) 63:1912–7. doi: 10.1002/1097-0424(19890515)63:10<1912::AID-cnrc28206301007-3.0.co2-y
4. Salvadori B, Marubini E, Miceli R, Conti AR, Curri F, Sventun M, et al. Reoperation for Locally Recurrent Breast Cancer in Patients Previously Treated With Conservative Surgery. Br J Surg (1989) 86:84–7. doi: 10.1046/j.1365-2168.1999.00961.x
5. Fourquet A, Campaana F, Zafirani B, Bossiri V, Vichl P, Durand JC, et al. Prognostic Factors of Breast Recurrence in the Conservative Management of Early Breast Cancer: A 25-Year Follow-Up. Int J Radiat Oncol Biol Phys (1989) 17:719–25. doi: 10.1016/0360-3016(89)90057-6
6. Touboul E, Buffat L, Belkacemi Y, Lefranç P, Uzan S, Lhuillier P, et al. Local Recurrences and Distant Metastases After Breast-Conserving Surgery and Radiation Therapy for Early Breast Cancer. Int J Radiat Oncol Biol Phys (1999) 43:25–38. doi: 10.1016/S0360-3016(99)00365-4
7. Peto R, Davies C, Godwin J, Gray R, Pan HC, Clarke M, et al. Comparisons Between Different Polychemotherapy Regimens for Early Breast Cancer: Meta-Analyses of Long-Term Outcome Among 100,000 Women in 123 Randomised Trials. Lancet (2012) 379:432–44. doi: 10.1016/S0140-6736(11)61625-5
8. Dalberg K, Mattsson A, Sandelin K, Rutqvist LE. Outcome of Treatment for Ipsilateral Breast Tumor Recurrence in Early-Stage Breast Cancer. Breast Cancer Res Treat (1998) 49:69–78. doi: 10.1023/a:1005934513072
9. Vila J, Garcia-Etienne CA, Vavassori A, Gentili O. Conservative Surgery for Ipsilateral Breast Tumor Recurrence. J Surg Oncol (2014) 110:62–7. doi: 10.1002/jso.23629
10. Chen SL, Martinez SR. The Survival Impact of the Choice of Surgical Procedure After Ipsilateral Breast Cancer Recurrence. Am J Surg (2008) 196:495–9. doi: 10.1016/j.amjsurg.2008.06.018
11. Su Y, Guo R, Xue J, Chi Y, Chi W, Wang J, et al. Increased Mortality With Repeat Lumpectomy Alone After Ipsilateral Breast Tumor Recurrence. Oncologist (2019) 24:e818–818e827. doi: 10.1634/thecologist.2018-0606
12. Wapnir IL, Gelber S, Anderson SJ, Mamounas EP, Robidoux A, Martin M, et al. Poor Prognosis After Second Locoregional Recurrences in the CALOR Trial. Ann Surg Oncol (2017) 24:398–406. doi: 10.1245/s10434-016-5571-y
13. Alpert TE, Kuerer HM, Arthur DW, Linnain DR, Haffy BG. Ipsilateral Breast Tumor Recurrence After Breast Conservation Therapy: Outcomes of Salvage Mastectomy vs. Salvage Breast-Conserving Surgery and Prognostic Factors for Salvage Breast Preservation. Int J Radiat Oncol Biol Phys (2005) 63:845–51. doi: 10.1016/j.ijrobp.2005.02.035
14. Fodor J, Major T, Polgar C, Orosz Z, Sulyok Z, Kasler M. Prognosis of Patients With Local Recurrence After Mastectomy or Conservative Surgery for Early-Stage Invasive Breast Cancer. Breast (2008) 17:302–8. doi: 10.1016/j.breast.2007.11.004
15. Komoki Y, Akiyama F, Iino Y, Ikeda T, Tanaka-Akashi S, Ohsumi S, et al. Analysis of Ipsilateral Breast Tumor Recurrences After Breast-Conserving Treatment Based on the Classification of True Recurrences and New Primary Tumors. Breast Cancer (2005) 12:104–11. doi: 10.2325/jbc.12.104
16. Lé MG, Arriagada R, Spielmann M, Guinebretière JM, Rocard F. Prognostic Factors for Death After an Isolated Local Recurrence in Patients With Early-Stage Breast Carcinoma. Cancer (2002) 94:2813–20. doi: 10.1002/cncr.10572
17. Solin LJ, Fourquet A, Vicini FA, Haffly B, Taylor M, McCormick B, et al. Salvage Treatment for Local Recurrence After Breast-Conserving Surgery and Radiation as Initial Treatment for Mammographically Detected Ductal Carcinoma in Situ of the Breast. Cancer (2001) 91:1090–7. doi: 10.1002/1097-0424(20010315)91:6<1090::AID-CNCR1104>3.0.CO;2-D
18. van Tienhoven G, Voogd AC, Petereit JL, Nielsen M, Andersen KW, Mignolet F, et al. Prognosis After Treatment for Loco-Regional Recurrence After Mastectomy or Breast Conserving Therapy in Two Randomised Trials (EORTC 10801 and DBCG-82tm). EORTC Breast Cancer Cooperative Group and the Danish Breast Cancer Cooperative Group. Eur J Cancer (1999) 35:32–8. doi: 10.1016/s0959-8049(98)00301-3
19. Yoshida A, Takahashi O, Okumura Y, Arima N, Nakatsukasa K, Tanabe M, et al. Prognosis After Mastectomy Versus Repeat Lumpectomy in Patients With Ipsilateral Breast Cancer Recurrence: A Propensity Score Analysis. Eur J Oncol (2016) 42:474–80. doi: 10.1016/j.ejso.2016.01.011
20. Abner AL, Recht A, Eberlein T, Come S, Shulman L, Hayes D, et al. Prognosis Following Salvage Mastectomy for Recurrence in the Breast After Conservative Surgery and Radiation Therapy for Early-Stage Breast Cancer. J Clin Oncol (1993) 11:44–8. doi: 10.1200/JCO.1993.11.1.44
21. Kurtz JM, Amalric R, Brandone H, Ayme Y, Spitalier J-M. Results of Salvage Surgery for Mammary Recurrence Following Breast-Conserving Therapy. Ann Surg (1988) 207:347–51. doi: 10.1097/00000658-198803000-00021
22. Kurtz JM, Spitalier JM, Amalric R, Brandone H, Ayme Y, Jacquierme J, et al. The Prognostic Significance of Late Local Recurrence After Breast-Conserving Therapy. Int J Radiat Oncol Biol Phys (1999) 46:1887–93. doi: 10.1016/S0360-3016(99)00271-k
23. Lee JH, Lee SK, Park SM, Ryu JM, Paik HJ, Yi HW, et al. Independent Prognostic Factors for Overall Survival After Salvage Operation for Ipsilateral Breast Tumor Recurrence Following Breast-Conserving Surgery. J Breast Cancer (2015) 18:386–93. doi: 10.4048/jbcs.2015.18.4.386
24. McCreary DR, Chapman JA, Wall JL, Lickley LA. Characteristics of Local Recurrence Following Lumpectomy for Breast Cancer. Cancer Invest (1994) 12:568–73. doi: 10.3109/07357909409023041

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SUPPLEMENTARY MATERIAL
The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fonc.2021.734719/full#supplementary-material
25. Sellam Y, Shahadi ID, Gelernter I, Zippel D, Sklar-Leyv M, Symon Z, et al. Local Recurrence of Breast Cancer: Salvage Lumpectomy as an Option for Local Treatment. Breast J (2019) 25:619–24. doi: 10.1111/tbj.13290

26. Smáňko V, Mízziaros N, Ujhelyi M, Fröhlich G, Stelzer G, Major T, et al. Second Breast-Conserving Surgery and Interstitial Brachytherapy vs. Salvage Mastectomy for the Treatment of Local Recurrences: 5-Year Results. Brachytherapy (2019) 18:411–9. doi: 10.1016/j.brachy.2019.02.004

27. Vooogd AC, van Tienhoven G, Peterse HL, Crommelin MA, Rutgers EJ, van de Velde CJ. Local Recurrence After Breast Conservation Therapy for Early Stage Breast Carcinoma: Detection, Treatment, and Outcome in 266 Patients. Dutch Study Group on Local recurrence After Breast Conservation (BORST). Cancer (1999) 85:437–46. doi: 10.1002/(sici)1097-0142(19990115)85:2<437::aid-cncr23>3.0.co;2-1

28. Komoike Y, Motomura K, Inaji H, Kasugai T, Koyama H. Repeat Lumpectomy for Patients With Ipsilateral Breast Tumor Recurrence After Breast-Conserving Surgery. Preliminary Results. Oncology (2003) 64:1–6. doi: 10.1159/000066512

29. Gentilini O, Botteri E, Veronesi P, Sangalli C, Del CA, Ballardini B, et al. Effectiveness of Breast-Conserving Surgery and 3-Dimensional Conformal Partial Breast Reirradiation for Recurrence of Breast Cancer in the Ipsilateral Breast: The NRG Oncology/RT0G 1014 Phase 2 Clinical Trial. JAMA Oncol (2020) 6:75–82. doi: 10.1001/jamaoncol.2019.4320

30. Deutsch M. Repeat High-Dose External Beam Irradiation for in-Breast Tumor Recurrence After Previous Lumpectomy and Whole Breast Irradiation. Int J Radiat Oncol Biol Phys (2002) 53:687–91. doi: 10.1016/s1046-1271(02)02785-2

31. Kauer-Dorner D, Tello JI, Zanon G, Henriquez I, Finestres F, et al. Exeresis and Brachytherapy as Salvage Treatment for Local Recurrence After Conservative Treatment for Breast Cancer: Results of a Ten-Year Pilot Study. Int J Radiat Oncol Biol Phys (2010) 78:804–10. doi: 10.1016/j.ijrobp.2009.08.009

32. Veronesi U, Marubini E, Andreola S, Greco M, et al. Local Recurrences and Distant Metastases After Conservative Breast Cancer Treatments: Partly Independent Events. J Natl Cancer Inst (1995) 87:19–27. doi: 10.1093/jnci/87.1.19

33. Harms W, Budach W, Dunst J, Feyer P, Fietkau R, Haase W, et al. DEGRO Practical Guidelines for Radiotherapy of Breast Cancer VI: Therapy of Locoregional Breast Cancer Recurrences. Strahlenther Onkol (2016) 192:199–208. doi: 10.1007/s00066-015-0939-7

34. Smith TE, Lee D, Turner BC, Carter D, Haffty BG. True Recurrence vs. New Primary Ipsilateral Breast Tumor Relapse: An Analysis of Clinical and Pathologic Differences and Their Implications in Natural History, Prognoses, and Therapeutic Management. Int J Radiat Oncol Biol Phys (2000) 48:1281–9. doi: 10.1016/s0360-3016(00)01378-x

35. Janni W, Shabani N, Dimpll T, Starflinger I, Rjosk D, Peschers U, et al. Matched Pair Analysis of Survival After Chest-Wall Recurrence Compared to Mammary Recurrence: A Long-Term Follow Up. J Cancer Res Clin Oncol (2001) 127:455–62. doi: 10.1007/s004320000238

36. Whelan T, Clark R, Roberts R, Levine M, Foster G. Ipsilateral Breast Tumor Recurrence Postlumpectomy is Predictive of Subsequent Mortality: Results From a Randomized Trial. Investigators of the Ontario Clinical Oncology Group. Int J Radiat Oncol Biol Phys (1994) 30:11–6. doi: 10.1016/0360-3016(94)90513-4

37. Ashi S, Gelber S, Anderson SJ, Läng I, Robidoux A, Martin M, et al. Chemotherapy for Isolated Locoregional Recurrence of Breast Cancer (CALOR): A Randomised Trial. Lancet Oncol (2014) 15:156–63. doi: 10.1016/s1470-2241(13)70589-8

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