A meta-analysis of clinical trials assessing the effect of radiofrequency ablation for breast cancer

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Background: Radiofrequency ablation (RFA) is a minimally invasive thermal ablation technique. We conducted a meta-analysis based on eligible studies to assess the efficacy and safety of RFA for treating patients with breast cancer.

Methods: A literature search was conducted in PubMed, Embase, and Web of Science databases. Eligible studies were clinical trials that assessed RFA in patients with breast cancer. The outcomes included complete ablation rate, recurrence rate, excellent or good cosmetic rates, and complication rate. A random-effects or fixed-effects model was used to pool the estimate, according to the heterogeneity among the included studies.

Results: Fifteen studies, with a total of 404 patients, were included in this meta-analysis. Pooled results showed that 89% (95% confidence interval: 85%–93%) of patients achieved a complete ablation after RFA treatment and 96% of patients reported a good-to-excellent cosmetic result. Although the pooled result for recurrence rate was 0, several cases of relapse were observed at different follow-up times. No RFA-related complications were recorded, except for skin burn with an incidence of 4% (95% confidence interval: 1%–6%).

Conclusion: This meta-analysis showed that RFA can be a promising alternative option for treating breast cancer since it produces a higher complete ablation rate with a low complication rate. Further well-designed randomized controlled trials are needed to confirm the efficacy and safety of RFA for breast cancer.

Keywords: radiofrequency ablation, breast cancer, meta-analysis

Introduction

The treatment for early breast cancer has evolved over the past decade. Nowadays, there is an increasing demand for minimally invasive treatments for small breast cancer. Breast conservation therapy has largely replaced mastectomy for the surgical treatment of early-stage breast cancer, as several randomized trials have shown a similar survival outcome between them.1–3 Moreover, sentinel lymph node biopsy has been widely accepted as an effective method for evaluating axillary nodal status for node-negative breast cancer, which avoids unnecessary axillary lymph node dissection.4,5 Owing to these fundamental changes in the breast cancer treatment strategy, there is an impetus to replace the less-invasive surgery with ablation techniques in the treatment of primary tumor without surgery.

Several minimally invasive ablation techniques have been applied for the treatment of small breast cancer, including cryosurgery, laser ablation, and thermoablation. Among them, radiofrequency ablation (RFA) is one of the most promising techniques. The treatment efficacy of RFA has been investigated in several studies, and results showed that RFA would be most effective for the treatment of tumors <3 cm.6–8 The purpose of this meta-analysis is to assess the feasibility and safety of RFA in the treatment of patients with breast cancer.
Materials and methods

Literature search

We conducted a comprehensive literature search in PubMed, Embase, and Web of Science databases from their inception through August 18, 2015. The following search terms were used: (“breast neoplasms”[MeSH Terms]) OR (“breast”[All Fields] AND “neoplasms”[All Fields]) OR (“breast neoplasms”[All Fields]) OR (“breast”[All Fields] AND “cancer”[All Fields]) OR (“breast cancer”[All Fields]) AND (radiofrequency [All Fields] AND ablation [All Fields]). The search was limited to human subjects, and no language restriction was imposed. We also manually searched the reference lists of included trials and reviews until no potential eligible trials could be found.

Review strategy

We used Endnote bibliographic software to build up an electronic library of citations identified in the literature searches. PubMed, Embase, and Web of Science databases were searched using Endnote, and duplicate records were deleted. Two independent investigators (CJ and ZC) were trained to perform the abstract/title review and then full-text review. Discrepancies between the investigators were resolved by consensus and discussion.

Study inclusion and exclusion criteria

All clinical trials assessing the efficacy and safety of RFA as a minimally invasive strategy for breast cancers were considered eligible for analysis. The following inclusion selection criteria were applied: 1) study population: adult women older than 18 years and had mammographic or ultrasonographic or magnetic resonance imaging evidence of breast cancer; 2) intervention: RFA; and 3) outcome measure: complete ablation rate and complication. Review articles, abstracts, editorials, case reports, and letters were not included in this meta-analysis.

Data extraction

Two independent investors (CJ and LF) extracted the following data from the included studies: first author, year of publication, number of patients, baseline character of patients, complete ablation rate, recurrence rate, excellent or good cosmetic rate, and incidence of complications. A standardized Excel file (Microsoft Corporation, Redmond, WA, USA) was established to extract the data from each study. When several publications from the same trial appeared, we included only the most informative article or the latest study to avoid duplication of information.

Statistical analysis

Before the data were pooled, we tested the heterogeneity among the studies using $F$ statistics. The studies were considered to have low, moderate, or high heterogeneity when the value of $F$ was 25%–50%, 50%–75%, or >75%, respectively. A value of $F$>50% indicates significant heterogeneity. When heterogeneity was found among the included studies, a random-effect model was used to pool the estimates; otherwise, a fixed-effect model was applied. Publication bias was evaluated using the test proposed by Begg and Mazumdar. A $P$-value of <0.05 was considered statistically significant. All analyses were performed using STATA Version 12.0 (StataCorp LP, College Station, TX, USA).

Results

Study identification and selection

The initial search yielded 1,678 records, of which 976 were excluded for the duplicate studies and 656 studies were excluded after a review of title/abstract, and the remaining 46 potential studies were included for the full-text review. Of the 46 studies for final analysis, 31 were excluded for the following reasons: 18 were unrelated with our topics, five were animal experiment, three did not provide outcomes of interest, and five used other ablation rather than RFA method. Thus, 15 studies met the inclusion criteria, and they were included in this meta-analysis (Figure 1).

Characteristics of eligible studies

The baseline characteristics of the included studies are presented in Table 1. These studies were published between 2001 and 2012. The sample sizes in these studies ranged from 10 to 52 (total of 404). The age of the patients ranged from 33 to 89 years, with a mean age >50 years. Of the 15 studies, eight studies reported that the tumor size was <2 cm, five studies reported <3 cm, and the remaining two studies reported <5 cm. The greatest tumor size was 5 cm, which was reported by Tsuda et al. Among these studies that presented the tumor histology, invasive ductal carcinoma was the primary type of carcinoma, which accounted for 74.7%. All the studies used ultrasound as image guidance to perform RFA. The mean temperature and time for tumor ablation were 95°C and 15 minutes, except for the method used by Wiksell et al., in which the tissue temperature and time were applied at 85°C and 10 minutes.

Complete ablation rate

Eleven studies reported the data of complete ablation rate. The complete ablation rate among these
studies ranged from 76% to 96.15%. Pooled estimates suggested that among the patients receiving RFA, 89% (95% confidence interval [CI]: 85%–93%) achieved a complete ablation (Figure 2). No heterogeneity was identified across these included studies ($I^2=0.0\%$, $P=0.454$).

Excellent cosmetic and good cosmetic

Four studies provided the data of cosmetic results.\textsuperscript{19,23,26,27} The cosmetic was scored according to the Radiation Therapy Oncology Group cosmesis scoring criteria. The aggregated results of these studies suggested that, of the patients treated

### Table 1 Baseline characteristics of patients in the trials included in the meta-analysis

<table>
<thead>
<tr>
<th>Study</th>
<th>Number</th>
<th>Mean age (range, years)</th>
<th>Tumor size (cm)</th>
<th>Tumor histology</th>
<th>Follow-up (months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hayashi et al\textsuperscript{15}</td>
<td>22</td>
<td>73 (60–80)</td>
<td>$\leq 1.0$</td>
<td>NR</td>
<td>NR</td>
</tr>
<tr>
<td>Noguchi et al\textsuperscript{16}</td>
<td>10</td>
<td>54 (33–70)</td>
<td>$&lt;2.0$</td>
<td>Invasive ductal/intraductal carcinoma: 7/3</td>
<td>NR</td>
</tr>
<tr>
<td>Izzo et al\textsuperscript{17}</td>
<td>26</td>
<td>57 (37–78)</td>
<td>1.8 (0.7–3.0)</td>
<td>Infiltrating ductal/lobular/tubular: 17/6/3</td>
<td>NR</td>
</tr>
<tr>
<td>Wiksell et al\textsuperscript{18}</td>
<td>33</td>
<td>NR (46–83)</td>
<td>0.6–1.5</td>
<td>Ductal/lobular/mucinous/tubular: 26/2/1/4</td>
<td>55±21</td>
</tr>
<tr>
<td>Wilson et al\textsuperscript{19}</td>
<td>40</td>
<td>68.8±10.9</td>
<td>0.2–2.6</td>
<td>NR</td>
<td>1–60</td>
</tr>
<tr>
<td>Palussière et al\textsuperscript{20}</td>
<td>21</td>
<td>79 (70–88)</td>
<td>$&lt;3.0$</td>
<td>Invasive ductal/lobular/mucinous carcinoma: 17/3/1</td>
<td>49.6 (17–77)</td>
</tr>
<tr>
<td>Burak et al\textsuperscript{21}</td>
<td>10</td>
<td>53.7 (37–67)</td>
<td>1.15 (0.8–1.5)</td>
<td>NR</td>
<td>NR</td>
</tr>
<tr>
<td>Ohtani et al\textsuperscript{22}</td>
<td>41</td>
<td>59 (38–92)</td>
<td>$&lt;2.0$</td>
<td>Invasive ductal/noninvasive ductal carcinoma: 36/5</td>
<td>NR</td>
</tr>
<tr>
<td>Medina-Franco et al\textsuperscript{23}</td>
<td>25</td>
<td>55.3 (42–89)</td>
<td>2.08 (0.9–3.8)</td>
<td>Infiltrating ductal/lobular/mixed carcinoma: 21/2/2</td>
<td>1–6</td>
</tr>
<tr>
<td>Fornage et al\textsuperscript{24}</td>
<td>21</td>
<td>56 (38–80)</td>
<td>$&lt;2.0$</td>
<td>Invasive ductal/lobular carcinoma: 19/2</td>
<td>NR</td>
</tr>
<tr>
<td>Ehrashi et al\textsuperscript{25}</td>
<td>17</td>
<td>55 (33–78)</td>
<td>1.1 (0.5–2.4)</td>
<td>Invasive ductal/noninvasive ductal carcinoma: 21/3</td>
<td>1–6</td>
</tr>
<tr>
<td>Oura et al\textsuperscript{26}</td>
<td>52</td>
<td>55 (37–83)</td>
<td>1.3 (0.5–2.0)</td>
<td>Ductal/invasive ductal/lobular/tubular: 7/42/2/1</td>
<td>Every 2–3</td>
</tr>
<tr>
<td>Yamamoto et al\textsuperscript{27}</td>
<td>29</td>
<td>55.9 (38–78)</td>
<td>1.28 (0.5–1.9)</td>
<td>NR</td>
<td>17 (2–41)</td>
</tr>
<tr>
<td>Seki et al\textsuperscript{28}</td>
<td>15</td>
<td>61 (36–82)</td>
<td>1.7 (0.5–3.0)</td>
<td>Invasive ductal carcinoma: 15</td>
<td>18</td>
</tr>
<tr>
<td>Tsuda et al\textsuperscript{29}</td>
<td>28</td>
<td>36–82</td>
<td>2.21 (0.6–5.0)</td>
<td>Intraductal carcinoma: 28</td>
<td>NR</td>
</tr>
</tbody>
</table>

Abbreviation: NR, not reported.
with RFA, 77% (95% CI: 57%–97%) rated their cosmetic result as excellent and 19% (95% CI: 4%–34%) rated as good (Figure 3). Overall, 96% of patients reported a good-to-excellent cosmetic result.

Recurrent rate
Five studies presented the data of recurrent rate.\textsuperscript{19,20,25–27} Of them, three\textsuperscript{25–27} reported that no recurrence occurred at the final follow-up. The follow-up of these studies was 1–7, 6–30, and 2–41 months, respectively. Whereas, in another study,\textsuperscript{20} one of the 21 patients reported a local relapse at the 1-year follow-up, and three additional patients presented with cancer recurrence outside the ablation zone at 30, 48, and 60 months. Moreover, the remaining study reported that one (1.3%) of 73 patients developed a true in-site recurrence at the 55±21 months follow-up.\textsuperscript{19} The pooled results indicated that patients treated with RFA had no local recurrence at a maximum follow-up of 76 months (Figure 4).

Complications
Seven studies reported the complications.\textsuperscript{15,17,20,22,23,26,27} No RFA-related complications were recorded by them. However, skin burn was noticed among these studies. Pooled estimates showed that the incidence of skin burn was 4% (95% CI: 1%–6%; Figure 5). And there was no significant heterogeneity among the studies ($I^2=18.9\%$, $P=0.286$).

Publication bias
The test proposed by Begg and Mazumdar\textsuperscript{14} was used to evaluate publication bias, and the result revealed that no potential publication bias existed among the included studies ($Z=0.78$, $P=0.436$).

Discussion
This study was a meta-analysis with the objective of assessing the efficacy and safety of RFA in the treatment of breast cancer. Our study indicated that among the patients with breast cancer who underwent RFA, 89% achieved a complete ablation, 96% reported a good-to-excellent cosmetic result, and 4% developed skin burn. This study indicated RFA is promising as a minimally invasive ablation technique in the local treatment of breast cancer.

RFA is a thermal ablation technique that has been applied for benign and malignant tumors.\textsuperscript{30,31} RFA causes local tumor cell death by thermal coagulation and protein denaturation.\textsuperscript{21,32,33} High-frequency alternating current flows from the tip of the electrode into the surrounding tissue causing ions to oscillate back and forth, thus generating frictional heat.\textsuperscript{34,35} Cell death approximately occurs $>45^\circ\text{C}$–$50^\circ\text{C}$.\textsuperscript{36} The targeted
temperature mostly used at the tip of prongs was 95°C and lasted for 15 minutes.\textsuperscript{15,24} This setting could melt the fatty tissue and also lead to bad cosmetic results. However, the lesions would achieve an equal ablation effect when RFA was set at a lower target temperature and a shorter ablation time.\textsuperscript{24} In this meta-analysis, we found that the complete ablation rate was 89%, indicating a high rate of complete tumor ablation for patients receiving RFA. The complete ablation rate among these studies ranged from 76% to 96.15%. The highest complete ablation rate was observed in the study...
Figure 5 Forest plot showing the complication rate of radiofrequency ablation in the treatment of patients with breast cancer. 
Abbreviations: CI, confidence interval; ES, estimates.

conducted by Izzo et al., in which ultrasound-guided RFA was performed in 26 patients with T1 and T2 breast tumors and then followed by immediate resection. Among these patients, 25 (96.15%) achieved a complete coagulation necrosis of tumor, while the remaining one had a microscopic focus of viable tissue adjacent to the needle shaft site. Another study performed RFA to treat patients with invasive breast carcinomas; however, the complete ablation rate was only 76%. In that study, the authors restricted patients to those with invasive breast cancers, no multicentric tumors, and no previous chemotherapy. Under ultrasound guidance, saline-cooled RFA was performed on tumors and a 5-mm margin of surrounding breast tissue, and then surgical resection was followed. Of the 25 patients, only 19 (76%) patients showed no evidence of viable malignant cells by the pathologic analysis of NDPH stain. The complete ablation rate for tumors \(< 2 \text{ cm} \) (92.8%) was significantly higher than that of tumors \( \geq 2 \text{ cm} \) (54.5%).

Regarding the recurrence, several studies revealed that no patients developed breast recurrence at their final follow-up, while the remaining studies reported a recurrence rate ranging from 1.37% to 14.29%. In the trial conducted by Oura et al., 52 patients with breast cancer received one session of RFA, for a maximum time of 30 minutes for the first 29 patients and 15 minutes for the following 23 patients. And the authors reported that no recurrence developed at the average follow-up of 15 months (6–30 months) after RFA. Similarly, in another trial conducted by Yamamoto et al., none of the 29 patients who underwent RFA recurred after a median follow-up of 17 months (2–41 months). However, in the trial conducted by Palussière et al., four patients developed cancer recurrence after RFA treatment. In that study, patients were all older than 70 years (70–88 years) and had undergone neoadjuvant endocrine therapy within the past 6 months. The tumor size was \(< 3 \text{ cm} \), and three patients were diagnosed with lobular carcinoma. During the 9-month follow-up, one patient was presented with a local relapse, and three additional patients developed cancer recurrence outside the ablation zone at 30, 48, and 60 months, two of whom were lobular carcinoma. Then, the authors suggest that RFA should not be recommended for lobular carcinoma.

With advances in diagnostic modalities, such as mammography, ultrasonography, and magnetic resonance imaging, the breast tumors with a diameter of \(< 1 \text{ cm} \) could be detected. Therefore, surgical treatment for breast cancer has become less invasive. However, the cosmetic problems still remain. In this study, we found that 96% of patients who underwent RFA reported a good-to-excellent cosmetic result. The result of RFA was higher than those of external beam radiation and partial breast irradiation, in which the rate of excellent-to-good cosmetic results ranged from 77.6% to 83.3%.
In terms of complications of RFA, there were seven studies reporting no RFA-related complications except for skin burn. Skin burn is the common complication of RFA in the treatment of breast cancer. The skin or underlying muscle close to the tumor is of concern because RFA can result in skin necrosis or chest wall burn. In the study conducted by Palussière et al., four of the 21 patients developed skin burn, which finally showed spontaneous healing after a maximum of 2 months. In another study conducted by Medina-Franco et al., in three of the 25 patients (12%) superficial skin burns occurred that required local excision. The authors pointed that there was no acute skin necrosis; however, they performed the surgical excision immediately after RFA treatment. Thus, the skin necrosis may not be well assessed since a delay in time period between RFA and resection would be necessary.

Although this meta-analysis indicated that RFA produced a higher complete ablation rate with a low complication rate, there are several problems that remain to be resolved: 1) lack of ability to precisely determine tumor size, 2) determination of 100% tumor cell killing, 3) ability to follow local recurrence, and 4) cosmetic outcome. Further studies are needed to determine whether the use of RFA for small breast cancer can provide local control and survival rates equivalent to those of conventional breast conservative surgery.

There are some potential limitations in this meta-analysis that should be considered. First, our study was conducted on 15 trials, and most of them had relatively small sample size. Compared with larger trials, small trials are more likely to result in overestimates of the treatment effect. Thus, our conclusion should be interpreted with caution. Second, there were variations in patients’ characteristics among the included studies, including breast size, tumor location, composition of the breast tissue (fat content), and vascularity of the breast. These factors may increase the heterogeneity and have a potential on the pooled results.

**Conclusion**

In conclusion, this meta-analysis showed that RFA can be a promising alternative option for invasive breast cancer since it produces a higher complete ablation rate with a low complication rate. However, considering that this study was conducted on small and nonrandomized controlled trials, the efficacy and safety of RFA for breast cancer should be confirmed with large-scale, well-performed trials.

**Disclosure**

The authors report no conflicts of interest in this work.


