Efficacy and Safety of Percutaneous Microwave Ablation and Cementoplasty in the Treatment of Painful Spinal Metastases and Myeloma

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*AJNR Am J Neuroradiol* 2018, 39 (7) 1376-1383
doi: https://doi.org/10.3174/ajnr.A5680
http://www.ajnr.org/content/39/7/1376

This information is current as of September 15, 2023.
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ABSTRACT

BACKGROUND AND PURPOSE: Painful spinal metastases are a common cause of cancer-related morbidity. Percutaneous ablation presents an attractive minimally invasive alternative to conventional therapies. We performed a retrospective review of 69 patients with 102 painful spinal metastases undergoing microwave ablation and cementoplasty to determine the efficacy and safety of this treatment.

MATERIALS AND METHODS: Procedures were performed between January 2015 and October 2016 with the patient under general anesthesia using image guidance for 102 spinal metastases in 69 patients in the following areas: cervical (n = 2), thoracic (n = 50), lumbar (n = 34), and sacral (n = 16) spine. Tumor pathologies included the following: multiple myeloma (n = 10), breast (n = 27), lung (n = 12), thyroid (n = 6), prostate (n = 5), colon (n = 4), renal cell (n = 3), oral squamous cell (n = 1), and adenocarcinoma of unknown origin (n = 1). Procedural efficacy was determined using the visual analog scale measured preoperatively and at 2–4 weeks and 20–24 weeks postprocedure. Tumor locoregional control was assessed on follow-up cross-sectional imaging. Procedural complications were recorded to establish the safety profile.

RESULTS: The median ablation time was 4 minutes 30 seconds ± 7 seconds, and energy dose, 4.1 ± 1.6 kJ. Median visual analog scale scores were the following: 7.0 ± 1.8 preoperatively, 2 ± 1.6 at 2–4 weeks, and 2 ± 2.1 at 20–24 weeks. Eight patients died within 6 months following the procedure. Follow-up imaging in the surviving patients at 20–24 weeks demonstrated no locoregional progression in 59/61 patients. Two complications were documented (S1 nerve thermal injury and skin burn).

CONCLUSIONS: Microwave ablation is an effective and safe treatment technique for painful spinal metastases. Further studies may be helpful in determining the role of microwave ablation in locoregional control of metastases.

ABBREVIATIONS: MWA = microwave ablation; ODI = Oswestry Disability Index; Pre = preprocedure; RT = radiation therapy; VAS = visual analog scale

BACKGROUND

Painful spinal metastases are a common cause of cancer-related morbidity. Percutaneous ablation presents an attractive minimally invasive alternative to conventional therapies. We performed a retrospective review of 69 patients with 102 painful spinal metastases undergoing microwave ablation and cementoplasty to determine the efficacy and safety of this treatment.

MATERIALS AND METHODS

Procedures were performed between January 2015 and October 2016 with the patient under general anesthesia using image guidance for 102 spinal metastases in 69 patients in the following areas: cervical (n = 2), thoracic (n = 50), lumbar (n = 34), and sacral (n = 16) spine. Tumor pathologies included the following: multiple myeloma (n = 10), breast (n = 27), lung (n = 12), thyroid (n = 6), prostate (n = 5), colon (n = 4), renal cell (n = 3), oral squamous cell (n = 1), and adenocarcinoma of unknown origin (n = 1). Procedural efficacy was determined using the visual analog scale measured preoperatively and at 2–4 weeks and 20–24 weeks postprocedure. Tumor locoregional control was assessed on follow-up cross-sectional imaging. Procedural complications were recorded to establish the safety profile.

RESULTS

The median ablation time was 4 minutes 30 seconds ± 7 seconds, and energy dose, 4.1 ± 1.6 kJ. Median visual analog scale scores were the following: 7.0 ± 1.8 preoperatively, 2 ± 1.6 at 2–4 weeks, and 2 ± 2.1 at 20–24 weeks. Eight patients died within 6 months following the procedure. Follow-up imaging in the surviving patients at 20–24 weeks demonstrated no locoregional progression in 59/61 patients. Two complications were documented (S1 nerve thermal injury and skin burn).

CONCLUSIONS

Microwave ablation is an effective and safe treatment technique for painful spinal metastases. Further studies may be helpful in determining the role of microwave ablation in locoregional control of metastases.

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Management of metastatic spine disease requires multidisciplinary input. Treatment options include not only continually evolving medical therapy regimens, surgical techniques, and radiation technologies but also emerging minimally invasive interventions. Treatment recommendations should be based on collaborative recommendations of specialists as outlined by the Metastatic Spine Disease Multidisciplinary Working Group, which has provided a consensus document detailing the evidence supporting their algorithms.

Image-guided ablative therapies demonstrate potential advantages compared with surgery and radiation therapy, including reduced morbidity, lower procedural cost, suitability for real-time imaging guidance, the ability to perform therapy in an outpatient setting, synergy with other cancer treatments, repeatability, and short procedural time. Ablation (alone or in combination with cementoplasty) can achieve a number of objectives: palliative pain control, skeletal stabilization, and tumor control. In most cases, microwave ablation offers the advantages of short procedural time, minimal invasiveness, supportability of their algorithms.
cases, an improvement in quality of life and prevention of mor-
bidity would be the primary goals, rather than tumor control,
especially in sick patients with limited life expectancy and poor
functional status, who are often poor surgical candidates.

An ever-expanding array of percutaneous ablation technolo-
gies is being used in the treatment of painful neoplastic lesions.
These modalities include cryoablation, ethanol ablation, laser ab-
lation, MRI-guided focused sonography ablation, and radiofre-
cuency ablation.

Microwave ablation (MWA) offers several potential advan-
tages over other ablative modalities that may increase its effective-
ness in the treatment of tumors, including larger tumor ablation
volumes with a single probe, faster ablation times, improved con-
vection profile, and optimal heating of necrotic masses. MWA has
consistently higher intratumoral temperatures than other heat-
based therapies. Like cryoablation, MWA can be performed with
multiple applicators. However, MWA is a relatively new in-
novation, and literature demonstrating its clinical efficiency in
treating spinal tumors is relatively sparse, with only 2 retrospec-
tive studies published to date (describing 20 and 37 lesions in 17
and 35 patients, respectively). The authors therefore aimed to
present a larger retrospective series to demonstrate the efficacy
and safety of microwave ablation and cementoplasty.

In our practice, polymethylmethacrylate is administered fol-
lowing ablation of spinal tumors. This is to prevent pathologic
fractures (either because of metastatic disease directly or follow-
ing ablative therapy), which may be a significant source of further
pain from metastatic disease. In addition to alleviating pain, this
may be helpful in preventing associated morbidities such as pro-
gressive kyphosis and respiratory compromise or damage to ad-
jacent spinal nerve roots.

The authors report the results of a retrospective review to evaluate
the safety and efficacy of MWA and cementoplasty for treating pain-
ful spinal metastases. To our knowledge, this series represents the
largest study yet published in the literature and serves to confirm or
refute findings from previously published smaller series.

We hypothesized that MWA ablation and cementoplasty
would result in markedly reduced pain scores. We further hy-
pothesized that a substantial proportion of our patients would
demonstrate no evidence of disease progression on 6-month fol-
low-up cross-sectional imaging.

The purpose of this study was to determine whether treatment
of metastatic disease to the vertebral body with MWA and cemen-
toplasty would significantly decrease visual analog scale (VAS)
pain scores at 6 months. The secondary purpose of this study was
to determine disease progression in the treated vertebral body in
patients with oligometastatic disease at 6 months with cross-sec-
tional imaging.

MATERIALS AND METHODS

Full Johns Hopkins Hospital institutional review board approval
was obtained for retrospective analysis. Informed consent for the
study was waived and Health Insurance Portability and Account-
ability Act compliance was guaranteed. Medical records, includ-
ing demographics and pain scores, were obtained and reviewed.

Efficacy of treatment was defined by accomplishing primary
and secondary objectives. The primary objective was pain reduc-
tion after microwave ablation and cementoplasty. The secondary
objective was effective disease control, based on the absence of
locoregional disease progression on follow-up cross-sectional im-
aging evaluation. Complications were reviewed in this cohort of
patients to assess the safety of the procedure.

Patient Information

A search was performed using the keywords “microwave abla-
tion,” “cementation,” “cementoplasty,” “kyphoplasty,” and “ver-
tebroplasty” on all files in the Electronic Medical Record of our
institution from January 2015 to October 2016. On the basis of
this search and application of the inclusion and exclusion criteria
detailed below, appropriate records were accessed.

A total of 105 patients were found in the search of our institu-
tion. These patients had all been screened at the multidisciplinary
conference. After we applied the inclusion and exclusion criteria,
69 patients (36 men, 33 women) with spinal metastases treated
with microwave ablation and cementoplasty from January 2015
to October 2016 were included in this retrospective review. The
mean age of the patients was 56.

Inclusion Criteria

We included the following:
A) Patients with intractable pain (VAS > 4) from solitary or
multiple sites of disease that resulted in a return visit to the on-
cologist (intractable pain was defined as unrelenting pain despite
treatment with narcotic anesthesia).
B) Patients who either
1) Had persistent or recurrent pain after radiation therapy (RT),
2) Were not candidates for RT,
3) Had declined RT,
4) Had tumor considered percutaneously accessible by imaging.

Exclusion Criteria

We excluded the following:
A) Patients with an alternative explanation for pain based on
correlation between clinical presentation and spine MR imaging.
B) Concurrent systemic illness,
C) Tumors with retropulsion of the posterior cortices, epidu-
ral extension, and/or cord compression,
D) Patients who had tumors whose margins approximated
nerve roots or the spinal canal.

Twenty-eight patients who had undergone prior radiation
therapy with continued pain were included in the study.

All patients had prior cross-sectional imaging (CT, MR imag-
ing, and/or PET/CT). Sixty-nine patients had complete assess-
ment of their pain in the interventional radiology clinic of our
department using the visual analog scale pain score preprocedur-
ally and at 2–3 weeks (n = 69) and 20–24 weeks (n = 61) post-
procedure. In addition, 52 patients were assessed using the Os-
westry Disability Index (ODI) score chart before the procedure
and at similar intervals following it, with 17 patients refusing to
use the ODI preset questionnaire for ODI determination.

Patients had been deemed appropriate for image-guided abla-
tion and cementation in the multidisciplinary conference after
discussion and review of cross-sectional imaging, which included
a combination of CT, PET/CT, and MR imaging. All cross-sec-

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tional studies were evaluated by a noninterventionalist, board-certified, fellowship-trained neuroradiologist and were subsequently reviewed by a board-certified, fellowship-trained spine interventional radiologist at the multidisciplinary tumor board.

On the basis of cross-sectional CT imaging, lesions were classified into 65 lytic, 13 sclerotic, and 24 mixed lytic/sclerotic. The tumor area ranged between 0.75 to 15.0 cm² (mean, 3.6 ± 2.2 cm²) based on CT and MR imaging review by the interventional radiologist. Figure 1 demonstrates an example of a lytic osseous metastasis in a patient with metastatic breast carcinoma.

Procedural Details
Microwave ablation was performed using the AveCure microwave ablation system (MedWaves, San Diego, California) under CT (16-slice Somatom Sensation; Siemens, Erlangen, Germany) or fluoroscopic (Axiom Artist dBA Bi-Plane; Siemens) guidance. Most procedures were performed as outpatient procedures (n = 63; 91.3%) with 4-hour postprocedural observation. Patients whose lesions were within 5 mm of nerve roots (n = 6; 8.7%) were observed for 23 hours. All procedures were performed with the patient under general anesthesia and prone. Lesions were measured in 3 planes on cross-sectional imaging before the procedure. These measurements were then used as a basis for calculating the volume (and correlating energy deposition and time parameters) for ablation as detailed in Table 1. The cross-sectional imaging was used in procedural planning, including making note of the proximity of tumoral margins to critical anatomic landmarks.

Local anesthesia (lidocaine hydrochloride 1%) was administered with a 3.5-inch 22-g spinal needle used to infiltrate the periosteum to ease the postprocedural pain followed by soft-tissue blunt dissection. An 11-g IVAS access cannula (Stryker Neurovascular, Kalamazoo, Michigan) was inserted via a posterior or posterolateral approach (either transpedicular or transcostovertebral) under image guidance and was positioned at the proximal edge of the tumor. A 13-g hand drill was then coaxially inserted into the tumor and advanced to its anterior edge. Once this position was confirmed, core biopsies were obtained and sent to the laboratory.

Subsequently, a single 14- or 16-g AveCure microwave ablation antenna (MedWaves, 1.8-mm, 20-cm-long) was coaxially inserted into the tumor. Placement of the hand drill and the ablation antenna is demonstrated in Fig 2. The size of the ablation antenna was determined on the basis of the calculated target lesion size, with the ablation zone calculated as 2 mm beyond the actual size of the target lesion to achieve adequate control. To calculate the appropriate cutoff temperature and energy deposition, we took the following approach: anteroposterior, lateral, and transverse measurements of the probe position relative to the vertebral margins. These measurements were then used to determine the appropriately sized antenna. Energy in kilojoules and temperature in centigrade were subsequently calculated using the vendor’s standard algorithm. The team was then able to achieve these end points with our ablation.

A 13-g Troxev curved needle (Stryker) was used to access lesions along the posterior aspect of vertebral body; then, a 16-g 90-cm flexible AveCure microwave antenna (Medwaves) was advanced into an adequate position for ablation. An automated drill (11-g Arrow On-Control; Teleflex, Morrisville, North Carolina) and a biopsy cannula were used to access mixed and dense sclerotic lesions with subsequent placement of the microwave probe.

For fluoroscopically guided procedures in which there was a small ablation, conebeam CT (InSpace Dyna CT; Siemens) was performed to confirm the antenna position through the lesion. Where there were large lesions encompassing two-thirds of the vertebral body, a bipedicular approach was used with overlapping ablation zones. For CT-guided procedures, the antenna position was always confirmed under CT. The burn size was overestimated by only 2 mm in all directions in patients with oligometastatic disease (to achieve better disease control). However, the accuracy

FIG 1. Axial CT images with the patient supine demonstrate lytic osseous metastasis in the T11(A) and T12(B) vertebral bodies in a 60-year-old woman with metastatic breast carcinoma.

| Table 1: Tabular results of Mann-Whitney U paired tests between VAS and ODI data collected Pre and at 2–4 and 20–24 weeks, respectively |
|----------------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|
| Table Analyzed                   | Mann-Whitney Test of VAS Pre–2/4 weeks | Mann-Whitney Test of VAS Pre–20/24 weeks | Mann-Whitney Test of ODI Pre–2/4 weeks | Mann-Whitney Test of ODI Pre–20/24 weeks |
| Column B                         | VAS Pre                           | VAS 2–4 weeks                    | ODI Pre                           | ODI 2–4 weeks                     |
| Column A                         | VAS 2–4 weeks                     | VAS 20–24 weeks                  | ODI Pre                           | ODI 20–24 weeks                   |
| Mann-Whitney test                |                                   |                                   |                                  |                                  |
| P value                          | <.0001                            | <.0001                            | <.0001                            | <.0001                            |
| Sum of ranks in columns A and B  | 5526, 1978                        | 5478, 2025                        | 2665, 1252                        | 2672, 1244                        |
| Mann-Whitney U                   | 86.5                              | 134                               | 261.5                             | 254                               |
| Median of column A               | 7, n = 61                         | 7, n = 61                         | 46, n = 44                         | 46, n = 44                         |
| Median of column B               | 2, n = 61                         | 2, n = 61                         | 24, n = 44                         | 24, n = 44                         |
| Difference: actual               | −5                                | −5                                | −22                               | −22                               |
| Difference: Hodges-Lehmann       | −5 to −4                          | −5 to −4                          | −28 to −16                         | −28 to −16                         |
| Exact 95.01% CI of difference    |                                   |                                   |                                   |                                   |
of the gross extent of induced coagulation was limited by both spatial and contrast resolution to approximately 2–3 mm, and imaging findings are only a rough guide to the success of the ablation therapy because microscopic foci of residual disease cannot be expected to be identified with standard imaging.

Thermocouples were not placed in our series because lesions with a breach of the posterior cortex were excluded from the study, but whenever the ablation zone was close to the neural element (<5 mm), we used repeat short ablation cycles to control diffusion of the heat zone without decreasing the effectiveness of ablation, a technique used by the spine interventionalist. However, use of thermocouples and somatosensory evoked potentials/motor-evoked potentials monitoring is advised to avoid thermal injury to the spinal cord or spinal nerves in cases in which these structures are approximated. We used hydrodissection in 1 patient with tumor close to the S1 nerve root with instillation of saline around the nerve.

Vertebral cement augmentation was performed for all lytic and mixed osseous tumors (n = 89) after technically successful ablation, which was defined by the ability to place the microwave antenna successfully in the center of the lesion and achieve the target ablation power based on the size of the target lesion. No cementation was performed for purely sclerotic lesions. Patients were observed for 3–4 hours in the post anesthesia care unit for any immediate postprocedural complications. Patients were contacted 24–48 hours following their procedures to inquire about their well-being. Any patients who reported new symptoms were brought into the clinic and reviewed.

**Statistical Analysis**

Mann-Whitney U tests and graphs were constructed for both ODI and VAS data preprocedurally, against data at 2–4 weeks and 3–6 months, respectively. The data consisted of the following: 69 patients with VAS and 52 patients with ODI scores obtained preprocedurally, at 2–4 weeks, and 20–24 weeks. Ablation time, temperature, energy, and duration of exposure were included as predictors in all models. Marginal plots were then constructed from the mixed models to visually compare the effects of predictors on outcomes. All statistical analyses were completed with GraphPad Prism software 7.0 (GraphPad Software, San Diego, California).

**RESULTS**

Technically successful ablations were achieved in 100% of patients. Technical success was defined as achieving a temperature of ≥80°C, with deposition of calculated energy based on the size of the target lesion with minimal-to-no reverse power.

The median ablation time was 4:30 ± 0.07 minutes (range, 3:00–10:34 minutes). The median energy dose received was 3.9 ± 1.6 kJ, and power was 13.28 ± 3.75 W. The preprocedural median VAS score was 7.0 ± 1.8. The median postprocedural VAS scores were 2 ± 1.6 at 2–4 weeks and 2 ± 2.1 at 20–24 weeks. A 4-point reduction in the VAS score was considered effective.

Follow-up imaging with CT of the chest, abdomen, and pelvis (n = 34) and PET/CT imaging20 at 6 months (n = 61) demonstrated stable disease in 59/61 patients (96.7%). There were 2 procedure-related complications (2.9%) (S1 nerve thermal injury with the patient having severe pain radiating to the leg in an S1 distribution with three-fifths motor strength, and a skin burn over the target lesion). Eight patients died within 6 months of the procedure.

The median Oswestry Disability Index was 46 ± 17.9 preprocedurally and 24 ± 17.1 and 24 ± 18.8 at 2–4 weeks and 20–24 weeks postprocedurally, respectively. Box-and-whisker plots for VAS and ODI scores are shown in Fig 3.

A median difference of −22 ± 12.8 and −5 ± 2.0 ODI and VAS points, respectively, was observed in the interval between preprocedure (Pre) and 3–6 months postprocedure. Most of the treated patients reported a decreased dosage or frequency of narcotic analgesic use. Changes in the VAS score are demonstrated in Fig 4. Tabular results of Mann-Whitney U paired tests between VAS and ODI data collected preprocedurally and at 2–4 and 20–24 weeks are detailed in Table 1.

Locoregional control at 20–24 weeks was achieved in all patients in the subgroup in our series who had oligometastatic disease (n = 22). Locoregional control was defined as stable disease at the treated level using a combination of conventional CT, PET/CT, and spine MR imaging. For the patient to be deemed stable, the imaging study was required to demonstrate the ablation zone encompassing the targeted area of tumor with no evidence of disease progression at the treated site.

Descriptive statistics showing quartiles, median, mean, SD, standard error of the mean, confidence intervals, and mean ranks for each dataset are detailed in Table 2.

**FIG 2.** Intraprocedural anteroposterior (A) and lateral (B) fluoroscopic images of T11- and T12-level microwave ablation. A microwave ablation probe (arrow) and a manual drill (arrowhead) have been advanced through access cannulas into the anterior aspect of the T11 and T12 vertebral bodies, respectively. Anteroposterior (C) and lateral (D) fluoroscopic images of the T11 and T12 vertebral bodies post-microwave ablation and cement augmentation. The access cannulas have been retracted into the pedicles.
As overall survival of cancer patients improves and medical imaging becomes increasingly ubiquitous, the number of patients with osseous metastases continues to grow. Microwave ablation represents a relatively new technique in the minimally invasive armamentarium that will likely play an increasing role in the palliative and potentially curative treatment of patients with painful spinal metastases. With this in mind, we reviewed results of microwave ablation and cementoplasty at our institution in the context of the existing scientific literature.

Microwave ablation of tumor cells is primarily achieved using electromagnetic methods, resulting in tumor destruction using devices with frequencies ranging between 900 and 2500 MHz. Electromagnetic microwaves heat matter by agitating water molecules in the affected and surrounding tissue, producing friction and heat, which induce cellular death via coagulation necrosis. Microwave ablation is more effective in high-impedance tissues like bone because poor thermal conduction in bone is a limiting factor in radiofrequency ablation. Osseous relative permeability and low conduction help microwaves penetrate deeper and are more effective in thermal ablation compared with radiofrequency ablation.

Radiofrequency ablation, cryoablation, MRI-guided focused ultrasound, and microwave are all being used in clinical practice to treat patients with painful osseous lesions, particularly metastases, for both pain relief and as a useful adjunct therapy for local tumor control. Microwave ablation of tumor cells is primarily achieved using electromagnetic methods, resulting in tumor destruction using devices with frequencies ranging between 900 and 2500 MHz. Electromagnetic microwaves heat matter by agitating water molecules in the affected and surrounding tissue, producing friction and heat, which induce cellular death via coagulation necrosis. Microwave ablation is more effective in high-impedance tissues like bone because poor thermal conduction in bone is a limiting factor in radiofrequency ablation. Osseous relative permeability and low conduction help microwaves penetrate deeper and are more effective in thermal ablation compared with radiofrequency ablation.

Radiofrequency ablation, cryoablation, MRI-guided focused ultrasound, and microwave are all being used in clinical practice to treat patients with painful osseous lesions, particularly metastases, for both pain relief and as a useful adjunct therapy for local tumor control. The heating of tissues is typically much faster and more precise with microwave ablation and causes better and complete coagulation necrosis.

Clinical evidence for microwave ablation in the spine has been limited to some small series. In a retrospective study, Kastler et al reported successfully treating 20 spinal metastases (17 patients) with MWA (with cementoplasty in 9 cases). They reported pain relief in 16 of 17 patients, with significant pain reduction 1 day, 1 week, 1 month, 3 months, and 6 months after ablation without any major complications.

In a second retrospective study, Pusceddu et al reported MWA of 37 osseous metastases (35 patients), which included spinal lesions in 12 patients. Local tumor control was achieved in all patients, and significant pain palliation, 1 week, 1 month, and 6 months after ablation (and 1 year after ablation in surviving patients at that time point). They also reported improved walking ability and no major complications.

In a prospective study, Aubry et al reported MWA of 16 bone and soft-tissue lesions in 13 patients. They reported an average necrosis rate of 85% at the first follow-up, and an 80% overall success rate; half of these patients were considered complete responders. Local control of the disease persisted with time, though residual disease progressed slowly. Time to recurrence in this study was 7.2 months.

The results of the current study recapitulate the findings in these prior works, with immediate pain reduction obtained in 94% (65/69) of these procedures, maintained for 6 months.

**Primary Outcome: Pain Relief**

This series affirms that microwave ablation is a feasible and effective procedure for pain relief in patients with refractory painful spinal metastases. Immediate pain reduction was obtained in 94% (65/69) of these procedures and maintained for >6 months.

Reduction in pain from ablation procedures is thought to be attributable to a combination of proposed mechanisms: destruction of pain nerve fibers in the periosteum and bone cortex with reduced pain transmission; reduction in the size of the tumor burden and volume with reduced transmission of pain via the nerve endings; and decreased osteoclastic activity and coagulative necrosis of the tumor cells with a resultant decrease in the production of nerve-stimulating cytokines such as interleukins and α-tumor necrosis factor.

There is a significantly increased risk of pathologic fracture following radiation therapy, particularly following stereotactic beam radiation therapy or stereotactic radiosurgery, with a reported incidence of between 15% and 40%. Fractures most commonly occur several weeks after radiation and typically involve lytic lesions. Thus, prophylactic stabilization of these lesions before RT may be useful. Despite the current lack of level I evidence to support this practice, we commonly perform vertebral augmentation before RT to prevent fracture-associated morbidity at our institutions.
**Structural Stabilization**

Patients with widely metastatic disease are often systemically unwell and are unable to tolerate prolonged conservative management and all it entails. Consequently, these patients benefit greatly from early stabilization of fractures, with a lower requirement for bed rest, bracing, and anesthesia. Even patients with chronic pain from unhealed fractures lasting longer than 12 months can benefit. In patients with metastatic disease, surgical decompression has been shown to provide a reasonable long-term ambulatory benefit. Early evaluation for surgical stabilization is therefore recommended, particularly in patients who are young and/or highly functional and have a reasonable long-term prognosis. However, invasive surgery may be inappropriate for frail patients with complex tumors. In addition, prolonged postoperative recovery can delay systemic therapy.

In this group of patients, in whom surgical decompression or fixation or both are inappropriate, there is strong evidence that percutaneous vertebral augmentation can achieve pain relief and functional improvement in both osteoporotic and pathologic fractures.

Cementoplasty (vertebroplasty when performed in the spine and sacroplasty in the sacrum), the injection of polymethylmethacrylate cement, stabilizes axial-loading areas of bone. This stabilization is important because vertebral body collapse after tumor ablation alone has been shown to occur in up to 60% of patients in the first year following ablation when not combined with cementoplasty support. Cementoplasty alone does not provide sufficient strength across areas of bone that experience significant shear or torsional stress. Traditionally, screw fixation would be required to account for such forces. New image-guided percutaneous interventions are currently being developed to facilitate reinforcement in these areas, sometimes termed “osteosynthesis.”

Vertebral augmentation has been shown to have acceptably low complication rates, quicker recovery times than surgery, and minimal disruption to adjunctive therapies, including chemotherapy and radiation. Patients are typically able to bear weight within a few weeks.

![Graphs](image-url)  
**FIG 4.** Comparing the change in VAS scores from preprocedure to 20–24 weeks postprocedure with ablation energy.

![Table](table-url)  
**Table 2: Descriptive statistics showing quartiles, median, mean, SD, SEM, confidence intervals, and mean ranks for each dataset**

|                      | VAS Pre | VAS 2–4 Weeks | VAS 20–24 Weeks | ODI Pre | ODI 2–4 Weeks | ODI 20–24 Weeks |
|----------------------|---------|---------------|-----------------|---------|---------------|-----------------|
| No. of values        | 61      | 61            | 61              | 44      | 44            | 44              |
| Minimum              | 2       | 0             | 0               | 20      | 4             | 2               |
| 25% Percentile       | 6       | 1             | 2               | 36      | 16            | 18              |
| Median               | 7       | 2             | 2               | 46      | 24            | 24              |
| 75% Percentile       | 8       | 3             | 3               | 58      | 34            | 35.5            |
| Maximum              | 10      | 6             | 6               | 92      | 64            | 60              |
| Mean                 | 6.869   | 2.098         | 2.443           | 48.5    | 25.64         | 25.86           |
| SD                   | 1.737   | 1.338         | 1.522           | 16.37   | 13.78         | 13.18           |
| SEM                  | 0.2224  | 0.1713        | 0.1949          | 2.469   | 2.077         | 1.988           |
| Lower 95% CI         | 6.424   | 1.756         | 2.053           | 43.52   | 21.45         | 21.86           |
| Upper 95% CI         | 7.314   | 2.441         | 2.833           | 53.48   | 29.83         | 29.87           |
| Mean ranks           | 90.58   | 32.42         | 33.2            | 60.56   | 28.44         | 28.27           |

**Note:** SEM indicates standard error of the mean.
hours after the procedure and are usually able to engage in physical therapy the following day to minimize further deconditioning. Given the rapid recovery time and absence of large surgical wounds, adjunctive chemotherapy and RT can be rapidly pursued with minimal interruption. With the development of effective adjunctive therapies, tumor regression may often invoke local instability. Vertebral augmentation may therefore also be used to offer stability to vulnerable lesions.

Our findings lend credence to the idea that microwave ablation (in combination with cementoplasty) is effective in achieving effective and durable pain relief and demonstrates an excellent adverse effect profile. Patients who had tumors whose margins approximated nerve roots or the spinal canal were excluded from this study. When these patients are treated, somatosensory-evoked potentials/motor-evoked monitoring is advised to ensure safety.

**Secondary Outcome: Tumor Control**

Oligometastatic disease is a unique pathologic state in which patients have limited disseminated disease that may potentially be curable. This model is gaining increasing support within the oncology community and provides justification for treating these patients with an aggressive approach (surgery or embolization). A5,15 While many patients have isolated osseous metastases, A6 surgical resection of osseous lesions is uncommon, likely due to morbidity of the operation. Stereotactic radiation therapy for oligometastases is an active area of investigation, showing promising early results. A6–38 A few series of image-guided ablation of limited metastatic disease have recently been published, including both exclusively osseous metastases and mixed disease. A5,15,39–42

Limited series in the literature show promising local control rates with oligometastatic disease. A19,43–46 In 1 series, percutaneous thermal ablation of metastatic spine lesions could achieve a reasonable rate of curative therapy (up to 67% at 1 year). A21 This result was achieved in a select subgroup of small, solitary lesions within the vertebral body without significant cortical destruction or posterior element involvement.

Tumor areas in patients in this study are detailed in Table 3. There was a small subset of patients in this study with oligometastatic disease (20/67). Cross-sectional imaging at 20–24 weeks demonstrated no evidence of locoregional progression. Further long-term follow-up would be required to compare this cohort with those described in the literature.

Definitive criteria to select appropriate patients for ablation of oligometastases have not yet been established. Most of the published series include patients with ≤5 sites of metastases. Image-guided ablation is usually used in nonsurgical candidates or in those who develop metastases in a previously irradiated field.

**Limitations of the Study**

The study was performed retrospectively on the basis of data from a single institution. Thus, a higher level of evidence could be achieved by performing a prospective, multicenter trial. A definitive record of the patients’ medical anesthetics was not kept, which would have potentially provided additional qualification of the effectiveness of therapy. Tumor ablation zones were measured and recorded as areas (rather than volumes). Although this approach may be slightly less accurate, it reflects our current clinical practice and that of several centers with whom we collaborate.

**CONCLUSIONS**

Microwave ablation is a promising, safe, and effective treatment for osseous tumors, resulting in both a reduction in pain and a degree of locoregional control of the disease process.

**ACKNOWLEDGMENTS**

We are grateful to David Yousem for manuscript revision and mentoring.

Disclosures: Jennifer S. Barr—UNRELATED: Board Membership: Musculoskeletal Transplant Foundation, Comments: member of medical board of trustees; money for my services goes to my institution into a resident funding for books/activities. A*Money paid to the institution.

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