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Radiofrequency Ablation of Miscellaneous Solid Tumors Excluding Liver Tumors

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Related Policies (if applicable)
SUR709.029: Radiofrequency Ablation (RFA) of Primary or Metastatic Liver Tumors

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Carefully check state regulations and/or the member contract.

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Coverage

Osteolytic Bone Metastases

Radiofrequency ablation **may be considered medically necessary** to palliate pain in individuals with osteolytic bone metastases who have failed or are poor candidates for standard treatments such as radiation or opioids.

Osteoid Osteomas

Radiofrequency ablation **may be considered medically necessary** to treat osteoid osteomas that cannot be managed successfully with medical treatment.

Renal Cell Carcinoma

Radiofrequency ablation **may be considered medically necessary** to treat localized renal cell carcinoma that is **no more than 4 cm** in size when criteria 1 and 2 are met:

1. When it is necessary to preserve kidney function in individuals with significantly impaired renal function (i.e., the individual has 1 kidney or renal insufficiency defined by a glomerular filtration rate of $<60 \text{ mL/min/m}^2$);
2. When the standard surgical approach (i.e., resection of renal tissue) is likely to worsen existing kidney function substantially; OR when the individual is not considered a surgical candidate.

Non-Small-Cell Lung Cancer

Radiofrequency ablation **may be considered medically necessary** to treat an isolated peripheral non-small-cell lung cancer lesion that is **no more than 3 cm** in size when criteria 1 and 2 are met:

1. When surgical resection or radiotherapy with curative intent is considered appropriate based on stage of disease, however, medical comorbidity renders the individual unfit for those interventions;
2. When the tumor is located at least 1 cm from the trachea, main bronchi, esophagus, aorta, aortic arch branches, pulmonary artery, and the heart.

Nonpulmonary Tumor(s) Metastatic To The Lung

Radiofrequency ablation **may be considered medically necessary** to treat malignant nonpulmonary tumor(s) metastatic to the lung that are **no more than 3 cm** in size when criteria 1 and 2 are met (see the Policy Guidelines section for additional criteria):

1. When it is necessary to preserve lung function because surgical resection or radiotherapy is likely to worsen pulmonary status substantially; OR when the individual is not considered a surgical candidate;
2. When there is no evidence of extrapulmonary metastases; AND the tumor is located at least 1 cm from the trachea, main bronchi, esophagus, aorta, aortic arch branches, pulmonary artery, and the heart.

Radiofrequency ablation **is considered experimental, investigational and/or unproven** as a technique for ablation of:

- Breast tumors;
- Lung cancer not meeting the criteria above;
- Renal cell cancer not meeting the criteria above;
- Osteoid osteomas that can be managed with medical treatment;
- Painful bony metastases as initial treatment; and
- All other tumors outside the liver including, but not limited to, the head and neck, thyroid, pancreas, adrenal gland, ovary, and pelvic/abdominal metastases of unspecified origin.

Policy Guidelines

The following are additional criteria developed by clinical judgment or consensus and existing guidelines for the use of radiofrequency ablation to treat metastatic tumors to the lung:

- No more than 3 tumors per lung should be ablated;

- Tumors should be amenable to complete ablation; AND
- Twelve months should elapse before a repeat ablation is considered.

Description

In radiofrequency ablation (RFA), a probe is inserted into the center of a tumor; then, prong-shaped, non-insulated electrodes are projected into the tumor. Next, heat is generated locally by an alternating, high-frequency current that travels through the electrodes. The localized heat treats the tissue adjacent to the probe, resulting in a 3 cm to 5.5 cm sphere of dead tissue. The cells killed by RFA are not removed but are gradually replaced by fibrosis and scar tissue. If there is a local recurrence, it occurs at the edge and can sometimes be retreated. RFA may be performed percutaneously, laparoscopically, or as an open procedure.

Radiofrequency Ablation

Radiofrequency ablation was initially developed to treat inoperable tumors of the liver. Recently, studies have reported on the use of RFA to treat other tumors. For some of these, RFA is being investigated as an alternative to surgery for operable tumors. Well-established local or systemic treatment alternatives are available for each of these malignancies. The hypothesized advantages of RFA for these cancers include improved local control and those common to any minimally invasive procedure (e.g., preserving normal organ tissue, decreasing morbidity, decreasing length of hospitalization).

Goals of RFA may include:

1. Controlling local tumor growth and preventing recurrence;
2. Palliating symptoms; and
3. Extending survival duration for patients with certain tumors.

The effective volume of RFA depends on the frequency and duration of applied current, local tissue characteristics, and probe configuration (e.g., single vs. multiple tips). RFA can be performed as an open surgical procedure, laparoscopically or percutaneously, with ultrasound or computed tomography guidance.

Potential complications associated with RFA include those caused by heat damage to normal tissue adjacent to the tumor (e.g., intestinal damage during RFA of kidney), structural damage along the probe track (e.g., pneumothorax as a consequence of procedures on the lung), and secondary tumors (if cells seed during probe removal).

Regulatory Status

The U.S. Food and Drug Administration (FDA) issued a statement in September 2008, concerning the regulatory status of RFA. The FDA has cleared RFA devices for the general indication of soft tissue cutting, coagulation, and ablation by thermal coagulation necrosis. Under this general indication, RFA can be used to ablate tumors, including lung tumors. Some RFA devices have been cleared for additional specific treatment indications, including partial or

complete ablation of nonresectable liver lesions and palliation of pain associated with metastatic lesions involving bone. The FDA has not cleared any RFA devices for the specific treatment indication of partial or complete ablation of lung tumors, citing lack of sufficient clinical data to establish safety and effectiveness for this purpose. The FDA has received reports of death and serious injuries associated with the use of RFA devices in the treatment of lung tumors.

Rationale

Medical policies assess the clinical evidence to determine whether the use of technology improves the net health outcome. Broadly defined, health outcomes are the length of life, quality of life (QOL), and ability to function - including benefits and harms. Every clinical condition has specific outcomes that are important to patients and to managing the course of that condition. Validated outcome measures are necessary to ascertain whether a condition improves or worsens; and whether the magnitude of that change is clinically significant. The net health outcome is a balance of benefits and harms.

To assess whether the evidence is sufficient to draw conclusions about the net health outcome of technology, 2 domains are examined: the relevance, and quality and credibility. To be relevant, studies must represent 1 or more intended clinical use of the technology in the intended population and compare an effective and appropriate alternative at a comparable intensity. For some conditions, the alternative will be supportive care or surveillance. The quality and credibility of the evidence depend on study design and conduct, minimizing bias and confounding that can generate incorrect findings. The randomized controlled trial (RCT) is preferred to assess efficacy; however, in some circumstances, nonrandomized studies may be adequate. RCTs are rarely large enough or long enough to capture less common adverse events and long-term effects. Other types of studies can be used for these purposes and to assess generalizability to broader clinical populations and settings of clinical practice.

Osteolytic Bone Metastases

Clinical Context and Therapy Purpose

The purpose of radiofrequency ablation (RFA) in individuals who have painful osteolytic bone metastases who have failed or are poor candidates for standard treatments is to provide a treatment option that is an alternative to or an improvement on existing therapies.

The following PICO was used to select literature to inform this policy.

Populations

The relevant population of interest is individuals with painful osteolytic bone metastases who have failed or are poor candidates for standard treatments. After lung and liver, bone is the third most common metastatic site and is relatively frequent among individuals with primary malignancies of the breast, prostate, and lung. Bone metastases often cause osteolysis (bone breakdown), resulting in pain, fractures, decreased mobility, and reduced QOL.

Interventions

The therapy being considered is RFA.

Comparators

The following therapies and practices are options to manage painful osteolytic bone metastases: medical management (e.g., chemotherapy) and radiotherapy. External-beam radiotherapy often is the initial palliative therapy for osteolytic bone metastases. However, pain from bone metastases is refractory to radiotherapy in 20% to 30% of individuals, while recurrent pain at previously irradiated sites may be ineligible for additional radiation due to risks of normal tissue damage. Other alternatives include hormonal therapy, radiopharmaceuticals (e.g., strontium 89), and bisphosphonates. Less often, surgery or chemotherapy may be used for palliation, and intractable pain may require opioid medications.

Outcomes

The general outcomes of interest are overall survival (OS), reduction in pain and medication use, fractures, functional outcomes, and QOL.

Patients would be followed for several years given the impact of bone metastases on bone remodeling.

Study Selection Criteria

Methodologically credible studies were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.
- To assess long-term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Cohort Studies

Levy et al. (2020) conducted a global, multicenter, nonrandomized, prospective postmarketing study to evaluate the effectiveness of RFA in patients with painful osteolytic bone metastases. (1) Between October 2017 and March 2019, 134 ablations were performed in 100 patients (68% vs. 32% of the cohort had a single vs. multiple sites treated, respectively). The most common tumor location was thoracic (44%) followed by lumbar (33%). Patient outcomes including pain, pain interference, and QOL were collected. Forty percent of the cohort did not participate through the 6-month follow-up, with 2 additional discontinuations after 6 months. The most common reason for discontinuation was death (30 patients), which were all classified as related to the underlying malignancy. The primary endpoint evaluated was pain improvement, from baseline to 3 months. At baseline, the mean score for worst pain (measured by Brief Pain Inventory) for the entire cohort was 8.2. After RFA, worst pain significantly improved, with mean scores decreasing to 5.6, 4.7, 3.9, 3.7, and 3.5 at 3 days, 1 week, 1 month, 3 months, and

6 months, respectively ($p<.0001$ for all visits). Immediate improvement in pain (≥ 2 -point change in worst pain at the treatment site[s] 3 days after RFA) was achieved by 59% of patients. Four adverse events were reported, of which 2 resulted in hospitalization for pneumonia and respiratory failure, respectively.

Case Series

Goetz et al. (2004) reported on an international study conducted at 9 centers in which 43 patients with painful osteolytic bone metastases were treated palliatively with RFA. (2) The study's primary outcome measure was the Brief Pain Inventory-Short Form, a validated scale from 0 (no pain) to 10 (worst pain imaginable). Patient eligibility required baseline values of 4 or more from 2 or fewer painful sites. Thirty-nine (91%) of the patients had previously received opioids to control pain from the lesion(s) treated with RFA, and 32 (74%) had prior radiotherapy to the same lesion. The mean pain score at baseline was 7.9 (range, 4 to 10). At 4, 12, and 24 weeks after RFA, average pain scores decreased to 4.5, 3.0, and 1.4, respectively (all $p<.001$). Forty-one (95%) patients achieved clinically significant reductions in pain scores, prospectively defined as a decrease of 2 units from baseline. Investigators also reported statistically significant ($p=.01$) decreases in opioid use at weeks 8 (by 59%) and 12 (by 54%).

An earlier case series by Gronemeyer et al. (2002) showed that palliative RFA provided significant pain relief in 9 (90%) of 10 patients with unresectable, osteolytic spine metastases who had no other treatment options. (3) Pain was reduced by an average of 74%; back pain-related disability was reduced by an average of 27%. Neurologic function was preserved in 9 patients and improved in the other. In another small case series, Kojima et al. (2006) assessed 24 patients with painful metastatic bone tumors who experienced pain-alleviating effects with RFA, which is consistent with other evidence. (4)

Section Summary: Osteolytic Bone Metastases

A prospective cohort study and case series have shown clinically significant reductions in pain relief (defined as a decrease of 2 units from baseline on the Brief Pain Inventory scale) or reductions in opioid use following treatment with RFA of osteolytic pain metastases in patients with no or limited treatment options. A multicenter, prospective study reported significant reductions in pain through the 6-month follow-up period, with 59% of patients achieving immediate improvement in pain within 3 days of RFA.

Osteoid Osteomas

Clinical Context and Therapy Purpose

The purpose of RFA in individuals who have painful osteoid osteomas is to provide a treatment option that is an alternative to or an improvement on existing therapies.

The following PICO was used to select literature to inform this policy.

Populations

The relevant populations of interest are individuals with painful osteoid osteomas.

Osteomas are the most common benign bone tumor, comprising 10% to 20% of benign and 2% to 3% of all bone tumors. They are typically seen in children and young adults, with most diagnosed in patients between 5 and 20 years of age. Osteomas are most common in the lower extremity (usually the long bones, mainly the femur) and less common in the spine. These tumors typically have a characteristic clinical presentation and radiologic appearance, with pain that is usually continuous and worse at night and commonly relieved by aspirin or other nonsteroidal anti-inflammatory drugs (NSAIDs). The natural history of the osteoid osteoma varies based on location, and although they rarely exceed 1.5 cm in diameter, may produce bone widening and deformation, limb length inequality, or angular deviations when near a growth plate. When located in the spine, these lesions may lead to painful scoliosis or torticollis. Sometimes they heal spontaneously after 3 to 7 years.

Interventions

The therapy being considered is RFA.

RFA of osteoid osteoma is done with a needle puncture, so no incision or sutures are needed; further, patients may immediately walk on the treated extremity and return to daily activities when the anesthetic effect wears off. The risk of recurrence with RFA of an osteoma is 5% to 10%, and recurrent tumors can be retreated with RFA. In general, RFA is not performed in many spinal osteomas because of possible thermal-related nerve damage.

Comparators

The following therapies and practices are options to treat osteoid osteomas: medical management, surgical excision, core drill excision, and laser photocoagulation.

Treatment options include medical management with NSAIDs, surgical excision (wide/en bloc excision or curetting), or the use of computed tomography (CT) or magnetic resonance imaging (MRI)-guided minimally invasive procedures including core drill excision, laser photocoagulation, or RFA. For many years, complete surgical excision was the classic treatment of osteomas, usually performed in patients with pain despite medical management. However, a substantial incision may be necessary, with the removal of a considerable amount of bone (especially in the neck of the femur). This increases the need for bone grafting plus internal fixation (which often necessitates a second procedure to remove the metalwork). Other possible risks include avascular necrosis of the femoral head and postoperative pathologic fracture. In addition, surgical excision leads to a lengthier convalescence and postoperative immobilization. Anatomically inaccessible tumors may not be completely resectable and may recur.

Outcomes

The general outcomes of interest are reductions in pain and medication use, normal bone development, and postsurgical adverse events.

Patients would be followed through adolescence to ensure normal skeletal development.

Study Selection Criteria

Methodologically credible studies were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.
- To assess long-term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Systematic Reviews

Tordjman et al. (2020) conducted a systematic review on CT-guided RFA for osteoid osteomas. (5) The review included 69 studies (43 retrospective and 12 prospective studies; rest of study designs were not identifiable) comprising 3023 patients. The weighted overall failure rate was 8.3% for the entire cohort. When studies were analyzed by time period conducted, the failure rate was significantly lower in studies conducted between 2011 and 2019 compared to those conducted between 2002 and 2010 (7% vs. 14%, $p=.004$). The complication rate for the entire cohort was 3%, with skin burns (0.7%) and infections (0.5%) as the most commonly reported.

Lanza et al. (2014) reported on a systematic review of various ablative techniques for osteoid osteomas. (6) Included in the review were 23 articles on RFA, 3 on interstitial laser ablation, and 1 with a combination of ablation techniques, totaling 27 articles (N=1772). The mean technical success was 100% and clinical success, defined as being pain-free, ranged from 94% to 98%, depending on the length of follow-up. Complications occurred in 2% of patients and included skin or muscle burn in 9 patients, 4 infections, nerve lesions or tool breakage in 3 patients each, delayed skin healing, hematoma, and failure to reach target temperature in 2 patients each, and fracture, pulmonary aspiration, thrombophlebitis, and cardiac arrest in 1 patient each. Eighty-six patients had tumor recurrence.

Retrospective Studies

In their retrospective study of the efficacy and complications of CT-guided RFA of spinal osteoid osteoma, Albisinni et al. (2017) concluded that CT-guided RFA is as effective as first-line therapy for the disease. (7) After RFA, clinical symptoms were evaluated at 3, 6, and 12 months, with a final evaluation at the end of the study. Results showed complete regression of osteoid osteoma symptoms in 57 (93.4%) of 61 ($p=.001$) patients observed between 2002 and 2012. Study limitations included the retrospective design and focus on a single treatment.

Lassalle et al. (2017) conducted a single-center, retrospective analysis of long-term outcomes for CT-guided RFA in 126 patients with suspected osteoid osteoma. (8) The study was conducted from 2008 to 2015. Phone evaluations were performed. The overall success rate was 94.3% among the 88 patients who participated in the follow-up calls. The study was limited by its retrospective design, imprecision of patients' memory over follow-up, the lack of clinical and imaging follow-up, and an inability to perform multivariate statistical analysis of factors associated with treatment failure.

Rimondi et al. (2012) reported on a retrospective study of 557 patients treated with CT-guided RFA as primary treatment for nonspinal osteoid osteomas. (9) All patients were followed for a mean of 3.5 years (range, 0.5 to 9 years). Pain relief occurred in all 557 patients within the first week after RFA and continued in 533 (96%) patients who remained asymptomatic through their last follow-up. Pain recurrence occurred in 24 (4%) patients. Complications occurred in 5 patients and included thrombophlebitis, skin burn, broken electrode, and 2 procedures in which the RFA generator failed to reach maximum temperature.

Sahin et al. (2019) conducted a single-center, retrospective study that evaluated clinical pain symptoms to demonstrate the rapid relief of pain symptoms after CT-guided RFA for osteoid osteomas. (10) A total of 116 patients were included, and the efficacy success rate in the study was 98%. All patients reported immediate pain relief following the procedure, with scores of 0 or 1 on a 10-point visual analog pain scale within 24 hours. The mean duration of follow-up was 23 months, and pain relapse was reported in 2 of 108 patients available for follow-up. Seven minor complications were reported after the procedure with superficial skin burns as the most common complication (n=4).

Case Series

An observational study by Knudsen et al. (2015) evaluated long-term clinical outcomes after CT-guided RFA in patients diagnosed with osteoid osteoma located in the upper and lower extremities. (11) The study population included 52 patients with a typical clinical history and radiologically confirmed osteoid osteoma who received CT-guided RFA treatment from 1998 to 2014 at a Danish university hospital. The clinical outcome was evaluated based on patient-reported outcome measures and medical record review. The response rate was 52 (87%) of 60. After 1 RFA treatment, 46 (88%) of 52 patients experienced pain relief, and 51 (98%) of 52 patients had pain relief after repeat RFA. One patient underwent open resection after RFA. No major complications were reported; 4 patients reported minor complications including small skin burn, minor skin infection, and hypoesthesia at the needle entry point. In all, 50 (96%) of 52 patients were reported to be "very satisfied" with the RFA treatment.

Rosenthal et al. (2003) reported their experience over an 11-year period with 271 RFA procedures for osteoid osteomas in 263 patients. (12) The short-term outcome was evaluated to detect procedure-related problems; by this definition, all procedures were considered technically successful. Long-term clinical success data (defined as being free of pain without additional procedures) were available in 126 patients, with complete clinical success observed in 89%. For procedures performed as the initial treatment, the success rate was 91%.

Section Summary: Osteoid Osteomas

Numerous retrospective studies and case series, and systematic reviews of observational data have evaluated RFA for the treatment of painful osteoid osteomas. In a systematic review of thermal ablation techniques, clinical success (pain-free) was achieved in 94% to 98% of patients. Results have indicated that most patients (89% to 96%) remained pain-free at longer-

term follow-up. Another systematic review reported similar success rates noting an average 8.3% failure rate among patients receiving CT-guided RFA.

Localized Renal Cell Carcinoma

Clinical Context and Therapy Purpose

Radical nephrectomy remains the principal treatment of renal cell carcinoma (RCC); however, partial nephrectomy (PN) or nephron-sparing surgery has been shown to be as effective as radical nephrectomy, with comparable long-term recurrence-free survival rates, in a select group of individuals. Alternative therapy such as RFA is of interest in individuals with small renal tumors when preservation of renal function is necessary (e.g., in patients with marginal renal function, a solitary kidney, bilateral tumors) and in individuals with comorbidities that would render them unfit for surgery. Another consideration would be in individuals at high risk of developing additional renal cancers (e.g., von Hippel-Lindau disease).

The purpose of RFA in individuals who have localized RCC no more than 4 cm in size is to provide a treatment option that is an alternative to or an improvement on existing therapies.

The following PICO was used to select literature to inform this policy.

Populations

The relevant population of interest is individuals with localized RCC no more than 4 cm in size. Small renal masses, defined as 4 cm or less, are common findings on diagnostic imaging of the abdomen pelvis. Some of these masses are assessed to be suspicious for malignancy or have been identified by biopsy as a localized RCC. Tumors can be further categorized according to international tumor, nodes, metastasis (TNM) staging where cT1a is a clinically diagnosed tumor \leq 4 cm that is confined to the kidney without any nodal involvement.

Interventions

The therapy being considered is RFA.

Comparators

The following practice is currently being used to treat localized RCC: surgical excision; either total nephrectomy or PN.

Outcomes

The general outcomes of interest are recurrence rates and a reduction in rates of renal failure. Individuals should be followed for at least 10 years to monitor for tumor recurrence.

Study Selection Criteria

Methodologically credible studies were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.

- To assess long-term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Systematic Reviews

In a systematic review and meta-analysis, Huang et al. (2025) compared stereotactic body radiotherapy (SBRT) with RFA, microwave ablation, and cryoablation outcomes for primary localized renal cell carcinoma. (13) The review identified 133 studies that included 8,910 total patients, of whom 612 received SBRT, 2,503 received RFA, 2,069 received microwave ablation, and 3,726 received cryoablation. Overall, 1-, 2-, and 5-year local control rates were highest for SBRT (99%, 97%, and 95%) compared with RFA (96%, 95%, 92%), microwave ablation (97%, 95%, 86%), and cryoablation (95%, 94%, 90%). Subgroup analysis by tumor size showed comparable outcomes across modalities for tumors <4 cm, but for tumors ≥4 cm, SBRT achieved the highest local control at all time points. Regarding survival, cancer-specific survival at 1 year was 100% and consistent across all treatment groups. By 5 years, slight variations emerged, with SBRT showing the lowest rate at 95% (and RFA the highest at 100%). The heterogeneity across studies was moderate to high for most treatment modalities, which limits the strength of the conclusions.

In a systematic review and meta-analysis, Yanagisawa et al. (2022) compared differential clinical outcomes of patients treated with PN versus those treated with ablation techniques, including RFA, cryoablation, and microwave ablation, for cT1b and cT1a renal tumors. (14) They identified 27 studies with 13,996 total patients who received either PN or ablation for treatment of their tumors. Investigators found that in both cT1a and cT1b renal tumors, there were no differences in the percent decline of estimated glomerular filtration rates (eGFR) or in the overall complication rates between PN and ablation therapy. There was also no difference in cancer mortality rates between PN and ablation in patients with either cT1a or cT1b tumors. However, compared to ablation, PN was associated with a lower risk of local recurrence in patients with either tumor type. There was significant heterogeneity across studies, which limits conclusions.

In their systematic review and meta-analysis, Uhlig et al. (2019) compared oncologic, perioperative, and functional outcomes for PN with outcomes for various ablative techniques, including RFA and others, for small renal masses (mean diameter=2.53 to 2.84 cm). (15) They identified 47 moderate-quality studies, mostly retrospective, published from 2005 to 2017, with a total of 24,077 patients. Of these patients, 15,238 received PN and 1877 received RFA. The network meta-analysis used PN as the reference point. The overall results indicated that PN had better OS and local control over ablative techniques, but it was not significantly better for cancer-related mortality. In addition, ablation had fewer complications and better renal function outcomes. Across the studies included, patients treated by PN tended to be younger with less comorbidities compared with patients receiving thermal ablation—a consideration when assessing the outcomes for survival and local control.

In a systematic review and meta-analysis, Katsanos et al. (2014) reviewed 1 RCT and 5 cohort studies (N=587) assessing thermal ablation (RFA or microwave) or nephrectomy for small renal

tumors (size, 2.5 cm). (16) The local recurrence rate was 3.6% in both groups (relative risk, 0.92; 95% confidence interval [CI], 0.4 to 2.14; $p=.79$). Disease-free survival was also similar in both groups up to 5 years (hazard ratio [HR], 1.04; 95% CI, 0.48 to 2.24; $p=.92$). However, the overall complication rate was significantly lower in the patients undergoing ablation (7.4%) versus nephrectomy (11.1%; pooled relative risk, 0.55; 95% CI, 0.31 to 0.97; $p=.04$).

El Dib et al. (2012) conducted a meta-analysis evaluating RFA and cryoablation for small renal masses. (17) Selected were 11 RFA case series (426 patients) and 20 cryoablation case series (457 patients) published through January 2011. The mean tumor size was 2.7 cm (range, 2 to 4.3 cm) in the RFA group and 2.5 cm (range, 2 to 4.2 cm) in the cryoablation group. Mean follow-up times for the RFA and cryoablation groups were 18.1 and 17.9 months, respectively. Clinical efficacy, defined as cancer-specific survival rate, radiographic success, no evidence of local tumor progression, or distant metastases, did not differ significantly between groups. The pooled proportion of clinical efficacy for RFA was 90% (95% CI, 86% to 93%) and 89% (95% CI, 83% to 94%) for cryoablation.

Table 1 summarizes the characteristics of all systematic reviews. Table 2 contains the results of the largest and most recent of the reviews (Uhlig et al. [2019] and Yanagisawa et al. [2022]).

Table 1. Characteristics of Meta-Analyses Assessing Radiofrequency Ablation for Renal Masses

Study	Dates	Trials	Participants	N (Range)	Design	Duration
Huang et al. (2025) (13)	2000-2024	133	Patients who had received RFA, SBRT, CRA, or MWA for primary and localized renal cell carcinoma	8,910 (n/a)	Prospective, observational, RCTs	Median ranged from 24.5 to 28.7 mo
Yanagisawa et al. (2022) (14)	2005-2021	27	Patients who underwent PN, RFA, CRA, or MWA for small renal tumors (cT1a or cT1b)	13,996 (18-8818)	Prospective, retrospective, 1 RCT	14 mo to 6 y
Uhlig et al. (2019) (15)	2006-2017	47	Patients who had received PN, RFA, CRA, or MWA for small renal masses	24,077 (18-1803)	Prospective, retrospective, 1 RCT	3 to 82 mo
Katsanos et al. (2014) (16)	2007-2012	6	Patients with small renal tumors receiving RFA or nephrectomy	587 (69-150)	1 RCT, 5 cohort	Up to 6 y
El Dib et al. (2012) (17)	2000-2008	31	Patients who had received RFA or CRA	957 (n/a)	Case series	7 to 45.7 mo

		for renal tumors, regardless of size			
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CRA: cryoablation; mo: month(s); MWA: microwave ablation; n/a: data not available; PN: partial nephrectomy; RCT: randomized controlled trial; RFA: radiofrequency ablation; SBRT: stereotactic body radiotherapy; y: year(s).

The results table below does not include Katsanos et al. (2014) because of complete study overlap with Uhlig et al. (2019). El Dib et al. (2012) and Huang et al. (2025) are not included because the comparator in the studies selected were various ablative strategies, not surgery. Table A1 in the Appendix compares the trials included in these systematic reviews and meta-analyses.

Table 2. Results of Select Meta-Analyses Assessing Radiofrequency Ablation for Renal Masses

Study	Cancer-Specific Mortality, IRR		Local Recurrence, IRR		Complications, OR		Renal Function Decline, MD in eGFR	
	RFA	PN	RFA	PN	RFA	PN	RFA	PN
Uhlig et al. (2019) (15)	2.03	1.00	1.79	1.00	0.89	1.00	6.49	0.00
95% CI	0.81 to 5.08		1.16 to 2.76		0.59 to 1.33		2.87 to 10.10	
	RFA (events)	PN (events)	RFA (events)	PN (events)	RFA (events)	PN (events)	RFA (total decline)	PN (total decline)
Yanagisawa et al. (2022) (14)	cT1a: 27 cT1b: 8	cT1a: 113 cT1b: 18	cT1a: 64 cT1b: 32	cT1a: 59 cT1b: 34	cT1a: 126 cT1b: 50	cT1a: 204 cT1b: 62	cT1a: 176 cT1b: 154	cT1a: 217 cT1b: 184
Pooled RR (95% CI)	cT1a: 0.87 (0.57 to 1.31) cT1b: 0.80 (0.32 to 1.98)		cT1a: 0.43 (0.28 to 0.66) cT1b: 0.41 (0.23 to 0.75)		cT1a: 1.34 (0.90 to 2.00) cT1b: 1.08 (0.76 to 1.53)		cT1a: MD, 2.42 (-0.06 to 4.89) cT1b: MD, 0.73 (-3.76 to 5.23)	
I^2 (p-value)	cT1a: 0% (.62) cT1b: 0% (.76)		cT1a: 20% (.23) cT1b: 30% (.20)		cT1a: 63% (.003) cT1b: 22% (.26)		cT1a: 83% (.0004) cT1b: 0% (.71)	

CI: confidence interval; eGFR: estimated glomerular filtration rate; IRR: incidence rate ratio; MD: mean difference in glomerular filtration rate; OR: odds ratio; RFA: radiofrequency ablation; PN: partial nephrectomy; RR: risk ratio.

Randomized Controlled Trials

In an RCT, Liu et al. (2016) analyzed the safety and efficacy of the operative effects of percutaneous RFA in early-state RCC versus retroperitoneoscopic radical operation of RCC. (18) There were 35 women and 28 men included; race and ethnicity of participants were not described. The observation group was treated with percutaneous RFA and the control group with a radical retroperitoneoscopy. A total of 76 clinically confirmed diagnosed cases, from January 2011 to January 2013, with RCC, were randomized to the observation (n=41) or the control (n=35) groups. Operation time, blood loss during operation, length of stay, and incidence complications were lower in the control group ($p<.05$). Total efficacy, tumor-free

survival times, and survival rates did not differ statistically between groups ($p>.05$); however, percutaneous RFA reduced postoperative recovery time and was associated with fewer complications. Trial limitations included small sample size and the brief duration of follow-up.

Retrospective Studies

Relevant studies not included in the above systematic reviews and meta-analyses are described below.

Marshall et al. (2020) conducted a single-center, retrospective evaluation in 100 patients with 125 RCCs who received percutaneous RFA between 2004 and 2015. (19) Median follow-up in the study was 62.8 months. Five-year overall, cancer-specific, and local progression-free survival were 75%, 92%, and 92%, respectively. Ten-year overall, cancer-specific, and local progression-free survival were 32%, 86%, and 92%, respectively. The rate of local tumor progression was higher in patients with tumors >4 cm compared to those with tumors ≤ 4 cm, but the difference was not statistically significant (6% vs. 13%, $p=.466$). The study also noted no significant changes in eGFR from baseline to 2 to 3 years post-procedure (65.2 vs. 62.1 mL/min/1.73 m²; $p=.443$). The overall complication rate in the study was 9%. Limitations of the study include its retrospective design, lack of a control group, and selection bias where patients selected for RFA over surgical resection likely had worse baseline comorbidity status, which may have negatively impacted OS rates.

Dai et al. (2017) conducted a retrospective evaluation of 30 patients with 31 central renal tumors who underwent percutaneous RFA between 2005 and 2010 to assess the clinical efficacy and safety of image-guided percutaneous RFA of central RCC with adjunctive pyeloperfusion. (20) OS was 96.0% (95% CI, 88.4% to 100.0%) and progression-free survival at 5 years was 80.9% (95% CI, 65.8% to 95.9%). The investigators found that complications were significantly higher for tumors located within 5 mm of the renal pelvis or 0 mm of a major calyx (28.6% vs. 4.0%; $p<.05$) and major complications occurred in 5 (12.8%) of 39 RFA sessions. They concluded that image-guided percutaneous RFA combined with pyeloperfusion had satisfactory clinical efficacy in the treatment of renal tumor but may be associated with significant major complications. The retrospective design and the small sample base are limitations to this analysis.

Over 10 years, Dvorak et al. (2017) retrospectively evaluated the technical success as well as mid-term and long-term efficacy and safety of RFA and microwave ablation with guided CT in 64 patients with small, non-central renal tumors. (21) Ninety-one ablation procedures were performed on 68 tumors, 12 to 60 mm in size. Treatment was successful in 50 (73.5%) tumors; a second procedure was successful in 13 (19.1%) cases; and for the 5 largest tumors (range, 45 to 60 mm; 7.4%), a third treatment was required. Investigators concluded that percutaneous ablation is safe and effective in treating small, non-central renal tumors of the T1a group. The retrospective study design is the major limitation of this study.

A publication by Iannuccilli et al. (2016) reported a mean 34.1 month follow-up (range, 1 to 131 months) of RFA with intent to cure in 203 patients with renal tumors. (22) Patients referred for

RFA were at high risk or had refused surgery. Smaller tumors were treated with a single electrode with a 2 or 3 cm active tip. Larger tumors were treated with a cluster electrode with 3 active tips. Patients were assessed annually for the appearance of residual tumor at the treatment site, and 26 (13%) had residual disease. Treatment effectiveness was 87% during follow-up. The likelihood of recurrence was increased for tumors 3.5 cm or larger, clear cell subtype, and treatment temperature of 70° or less. All-cause mortality increased with increasing tumor size. The median survival was 7 years for patients with tumors less than 4 cm, with 80% survival at 5 years. Major complications, including urinary stricture or urine leak, occurred in 8 (3.9%) treatments.

Section Summary: Localized Renal Cell Carcinoma

The evidence on RFA for small renal tumors (≤ 4 cm) includes an RCT, meta-analyses, retrospective and cohort studies, and case series, that have compared RFA with nephrectomy or cryoablation. A 2014 meta-analysis that included 1 RCT and 5 cohort studies found that RFA was as effective as nephrectomy for small renal tumors, with a reduction in complications. Another, more recent, meta-analysis (2019) found that PN was superior to ablative techniques (the study included RFA but also cryoablation and microwave ablation) in overall mortality and local recurrence but not in cancer-specific mortality. It also found fewer complications and improved renal function with ablation. A meta-analysis from 2022 found that PN was superior to ablation (RFA, cryoablation, and microwave ablation) in local recurrence. Overall complications, decline in renal function, and cancer-specific mortality rates did not differ between ablation and PN. A 2025 meta-analysis found that RFA achieved high local control and cancer-specific survival rates comparable to other ablative therapies, although SBRT showed superior local control for larger tumors. The correlation between tumor size and RFA efficacy has been demonstrated by a large case series with a mean 34-month follow-up; it found that residual disease and mortality increased with tumors over 4 cm. Long-term follow-up in one single-center study found that RFA resulted in similar cancer-specific survival outcomes as PN in patients with cT1a renal tumors.

Primary Pulmonary and Nonpulmonary Tumors

Clinical Context and Therapy Purpose

Surgery is the current treatment of choice in individuals with stage I primary non-small-cell lung cancer (NSCLC; stage I includes 1a [T1N0M0] and 1b [T2N0M0]). Approximately 20% of patients present with stage I disease, although this number is expected to increase as a result of screening programs, advances in imaging modalities and widespread use of CT scans for other indications. Postsurgical recurrence rates of stage I NSCLC have been reported as between 20% and 30%, with most occurring at distant sites; locoregional recurrences occur in approximately 12%. Large differences in survival outcomes are observed after surgery in stage I disease, with 5-year OS rates ranging from 77% for small T1 tumors to 35% for large T2 tumors. Untreated, stage I NSCLC has a 5-year OS rate range from 6% to 14%.

Individuals with early-stage NSCLC who are not surgical candidates may be candidates for radiotherapy with curative intent. In 2 large, retrospective, radiotherapy series, patients with the inoperable disease treated with definitive radiotherapy achieved 5-year survival rates of

10% and 27%. In both studies, patients with T1N0 tumors had better 5-year survival rates of 60% and 32%, respectively.

Stereotactic body radiotherapy has gained more widespread use as a treatment option because it is a high-precision mode of therapy that delivers very high doses of radiation. Two- to 3-year local control rates of stage I NSCLC with stereotactic body radiotherapy have ranged from 80% to 95%. Stereotactic body radiotherapy has been investigated in individuals unfit to undergo surgery, with survival rates similar to surgical outcomes.

RFA also is being investigated in individuals with small primary lung cancers or lung metastases who are deemed medically inoperable.

The purpose of RFA in individuals who have inoperable primary pulmonary tumors or nonpulmonary tumors metastatic to the lung is to provide a treatment option that is an alternative to or an improvement on existing therapies.

The following PICO was used to select literature to inform this policy.

Populations

The relevant population of interest is individuals with inoperable primary pulmonary tumors or nonpulmonary tumors metastatic to the lung.

Interventions

The therapy being considered is RFA.

Comparators

The following practice is currently being used to treat primary pulmonary tumors or nonpulmonary tumors metastatic to the lung: radiotherapy.

Outcomes

The general outcomes of interest are OS, tumor recurrence, and treatment-related adverse events (e.g., pneumothorax). Individuals would be followed for at least 5 years.

Study Selection Criteria

Methodologically credible studies were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.
- To assess long-term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Systematic Reviews

In a systematic review of RFA, surgery, and stereotactic body radiotherapy for colorectal cancer (CRC) lung metastases, Schlijper et al. (2014) did not identify any randomized trials, and evidence was insufficient to draw conclusions on the comparative effectiveness of these therapies. (23)

In a comparative effectiveness review conducted for the Agency for Healthcare Research and Quality, Ratko et al. (2013) assessed local nonsurgical therapies for stage I NSCLC. (24) In this review, no comparative RFA studies were identified. Reviewers found that available evidence was insufficient to draw conclusions on the comparative effectiveness of local nonsurgical therapies for NSCLC, including RFA.

In a review of 16 studies, Bilal et al. (2012) compared RFA with stereotactic ablative radiotherapy in patients with inoperable early-stage NSCLC. (25) Reviewers found that OS rates for RFA and stereotactic ablative radiotherapy were similar in patients at 1 year (68.2% to 95% vs. 81% to 85.7%) and 3 years (36% to 87.5% vs. 42.7% to 56%), all respectively. However, survival rates at 5 years were lower with RFA (20.1% to 27%) than with stereotactic body radiotherapy (47%). These findings were drawn from comparisons of results from uncontrolled case series and retrospective reviews.

In an evidence-based review by Chan et al. (2011), 46 studies on RFA for lung tumors were evaluated, which included 2905 ablations in 1584 patients with a mean tumor size of 2.8 cm. (26) Twenty-four studies reported rates of local recurrence, which occurred in 282 (12.2%) cases at a mean follow-up of 13 months (range, 3 to 45 months). Primary lung cancer rates of local recurrence did not differ significantly (22.2%) from metastases (18.1%). Twenty-one studies reported mean OS rates of 59.4% at a mean follow-up of 17.7 months. The mean cancer-specific survival rate was 82.6%, at a mean follow-up of 17.4 months. The mean overall morbidity was 24.6% and most commonly included pneumothorax (28.3%), pleural effusion (14.8%), and pain (14.1%). Mortality related to the RFA procedure was 0.21%, overall.

Prospective Studies

A relevant study not included in the above systematic reviews and meta-analyses is described below.

Hasegawa et al. (2020) conducted a prospective, single-arm, multicenter study to evaluate the efficacy of RFA in patients with surgically resectable CRC lung metastases measuring 3 cm or smaller. (27) A total of 70 patients with CRC and 100 lung metastases were enrolled. All tumors were considered technically resectable, but not all patients were clinically able to undergo surgery. A total of 85 initial RFA sessions were performed for 100 target lung metastases. The 3-year OS rate after RFA was 84%. Primary and secondary technical success rates for RFA were 96% and 100%, respectively. Over a mean follow-up of 57 ± 32 months, local tumor progression was found in 6 patients (9%) at 6 to 19 months after the initial RFA. The 3-year progression-free survival rate was 41%. Grade 2 pneumothorax occurred after 18 of the 88 RFA sessions. The study is limited by its lack of a comparator arm.

Section Summary: Primary Pulmonary and Nonpulmonary Tumors

The evidence on RFA for primary NSCLC and nonpulmonary tumors metastatic to the lung includes prospective and observational studies and systematic reviews of those studies. No RCTs identified compared treatment approaches. For surgically resectable CRC lung metastases a multicenter study found that RFA for tumors less than 3 cm can lead to a 3-year OS rate of 84%. Two-year survival has been reported to range from 41% to 75% in case series. Survival at 1 and 2 years appears to be similar, following treatment with RFA or stereotactic ablative radiotherapy in patients with inoperable lung tumors. Survival rates at 5 years were lower with RFA (20.1% to 27%) than with stereotactic ablative radiotherapy (47%), but this finding was drawn from comparisons of uncontrolled case series and retrospective reviews. Prospective comparison in an RCT would permit greater certainty for this finding, but the studies are consistent with some effect of RFA on lung tumors.

Breast Tumors

Clinical Context and Therapy Purpose

The treatment of small cancers of the breast has evolved from total mastectomy to more conservative treatment options such as lumpectomy, with more acceptable cosmetic outcomes and preservation of the breast. The selection of the surgical approach balances the individual's desire for breast conservation and the need for tumor-free margins in resected tissue. Minimally invasive nonsurgical techniques such as RFA are appealing if they can produce local control and survival equivalent to breast-conserving surgical alternatives. Nonsurgical ablative techniques pose difficulties such as the inability to determine tumor size, complete tumor cell death, and local recurrence. Additionally, RFA can burn the skin and cause damage to muscle, possibly limiting use in patients with tumors near the skin or chest wall.

The purpose of RFA in individuals who have breast tumors is to provide a treatment option that is an alternative to or an improvement on existing therapies.

The following PICO was used to select literature to inform this policy.

Populations

The relevant population of interest is individuals with breast tumors.

Interventions

The therapy being considered is RFA.

Comparators

The following practices are currently being used as treatment options for small cancers of the breast cancer: radiotherapy and surgical excision.

Outcomes

The general outcomes of interest are tumor recurrence, reduction in medication, and treatment-related adverse events.

Patients would be followed for up to 5 years.

Study Selection Criteria

Methodologically credible studies were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.
- To assess long-term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Systematic Reviews

Xia et al. (2021) conducted a systematic review and meta-analysis of studies assessing RFA in patients with breast cancer and tumors that were 2 cm or smaller. (28) The primary endpoints of interest were technical success rate, complete ablation rate, and rate of complications. A total of 17 studies were identified, which accounted for 399 patients (401 lesions). Technical success rates ranged from 86.67% to 100% in the included studies; the pooled technical success rate was 99% (95% CI, 98% to 100%). After RFA, the majority of patients underwent surgical tumor excision (65.74%, 261/397). The pooled complete ablation rate was 98% (95% CI, 97% to 100%). The complication rate in the entire cohort was 6.8%; the most common complications were skin burns (2%), breast inflammation (1.5%), and infections (1%). The pooled complications rate was 2% (95% CI, 1% to 4%). Local recurrence was reported in 10 studies (232 cases); there was no local recurrence reported after a median follow-up of 27 months in these patients. The authors noted that prospective studies evaluating the use of RFA alone are needed to validate the place in therapy.

Peek et al. (2017) conducted a systematic review and meta-analysis of all studies evaluating the role of ablative techniques in the treatment of breast cancer published between 1994 and 2016. (29) Selection criteria included at least 10 patients with breast cancer treated with RFA, high-intensity ultrasound, or cryo-, laser, or microwave ablation; 63 studies (N=1608) were identified through PubMed and PubMed library databases. Fifty studies reported complete ablation, and RFA had the highest rate of complete ablation (87.1% [491/564]) as well as the shortest treatment time (15.6 minutes). A major limitation of this systematic review was the authors' inability to perform a comparative meta-analysis due to the inclusion of only 4 RCTs and 1 retrospective analysis that compared 2 or more techniques. There was also considerable heterogeneity across included studies.

Zhao and Wu (2010) conducted a systematic review of 38 studies on ablation techniques for breast cancer treatment published from 1994 to 2009. (30) Nine studies focused on RFA. Reviewers included small tumors ranging in size from 0.5 to 7 cm. Tumor resection was performed immediately after ablation or up to 4 weeks after RFA. Complete coagulation necrosis rates of 76% to 100% were reported. The results suggested RFA for breast cancer

tumors is feasible, but further studies with longer follow-up on survival, tumor recurrence, and cosmetic outcomes would be needed to establish clinical efficacy.

In another review, Soukup et al. (2010) examined 17 studies on RFA for the treatment of breast tumors and found RFA is feasible. (31) Even though few adverse events and complications occurred with breast RFA, incomplete tumor ablation remains a concern.

Clinical Studies

Relevant studies not included in the above systematic reviews and meta-analyses are described below.

Retrospectively, Ito et al. (2018) studied the safety and efficacy of percutaneous RFA of breast carcinomas in 386 patients from 10 institutions treated with RFA between 2003 and 2009. (32) Race and ethnicity of participants were not described. Patients were followed for a median of 50 months, and ipsilateral breast tumor recurrence was more frequent in patients with initial tumor sizes of 2 cm or more (10% [3/30]) than those with initial tumors 2 cm or less (2.3% [8/355]; $p=.015$). Ipsilateral breast tumor recurrence rates 5 years after RFA were 97%, 94%, and 87% in patients with initial tumor sizes of 1 cm or less, 1.1 to 2.0 cm, and greater than 2 cm, respectively. The authors concluded that RFA was safe for tumors of 2 cm or less. The retrospective design and lack of data on ipsilateral breast tumor recurrence for different types of chemotherapy and endocrine therapy and analyses to ascertain whether adjuvant chemotherapy or endocrine therapy influenced outcomes are the limitations of this study.

The efficacy and safety of using ultrasound-guided RFA for multiple breast fibroadenoma as an alternative to surgical resection were retrospectively analyzed by Li et al. (2016). (33) From 2014 to 2016, 65 patients with 256 nodules were treated with ultrasound-guided RFA, and complete ablation was achieved for 251 nodules (98.04%) after the first month of treatment; after the first and third months, tumor volume overall was reduced by 39.06% and 75.99%, respectively. The study reported minimal to no complications such as skin burns, hematoma, or nipple discharge. The retrospective design and short follow-up time limited the conclusions drawn from this study. Race and ethnicity of participants were also not described.

Wilson et al. (2012) reported on 73 patients with invasive breast cancer who had a lumpectomy followed immediately by RFA to the lumpectomy bed. (34) The average breast tumor size was 1.0 cm (range, 0.2 to 2.6 cm) and follow-up averaged 51 months. Disease-free survival was 100%, 92%, and 86% at 1, 3, and 5 years, respectively. One patient had tumor recurrence within 5 cm of the lumpectomy site, and 3 patients had ipsilateral breast recurrences.

Section Summary: Breast Tumors

Systematic reviews, retrospective studies, and observational studies have reported varied and incomplete ablation rates as well as concerns about post-ablation tumor cell viability. Long-term improvements in health outcomes have not been demonstrated. Additionally, available studies have not compared RFA with conventional breast-conserving procedures. For small breast tumors, further prospective study, with long-term follow-up, is needed to determine

whether RFA can provide local control and survival rates compared with conventional breast-conserving treatment.

Benign Thyroid Nodules

Clinical Context and Therapy Purpose

Surgical resection is the primary treatment choice for medically unresponsive, symptomatic benign thyroid tumors and thyroid carcinomas. However, techniques for ablation of thyroid tumors (e.g., RFA, microwave ablation) are being investigated.

The purpose of RFA in individuals who have benign thyroid tumors is to provide a treatment option that is an alternative to or an improvement on existing therapies.

The following PICO was used to select literature to inform this policy.

Populations

The relevant population of interest is individuals with large or symptomatic benign thyroid tumors. Individuals with a benign cytology diagnosis or those very unlikely to be malignant (e.g., purely cystic nodule) should undergo surveillance with the frequency determined by the level of suspicion for a missed malignancy. (35) Medical or surgical intervention is considered if the nodules are large (>4 cm), causing compressive or structural symptoms, or if there is clinical concern.

Interventions

The therapy being considered is RFA.

Comparators

The following practices are currently being used to treat large or symptomatic benign thyroid tumors in the United States: percutaneous ethanol injection (PEI) and surgical excision.

Outcomes

The general outcomes of interest are a reduction in nodule volume, hyper- and hypothyroidism, and treatment-related adverse events (e.g., voice changes).

Patients would be followed for at least 5 years.

Study Selection Criteria

Methodologically credible studies were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.
- To assess long-term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Systematic Reviews

Xu et al. (2024) evaluated the efficacy of thermal ablation (RFA, microwave, and laser ablation) for the treatment of benign thyroid nodules. (36) The analysis was limited to studies that had longer-term follow-up (approximately 5 years). A total of 5 studies (N=939) with 3 studies (n=483) specific to RFA were included. A total of 137 patients had local nodule recurrence at a median follow-up of 59.25 months.

Cho et al. (2020) evaluated the efficacy of thermal ablation (RFA and laser ablation) for the treatment of benign thyroid nodules. (37) The analysis demonstrated long-term maintenance (up to 36 months) of volume reduction. Further, RFA was found to be superior to laser ablation. The volume reduction rate for RFA at the last follow-up was 92.2%, whereas in the laser ablation group, the volume reduction rate peaked at 12 months (52.3%) and was at 43.3% at the last follow-up.

To evaluate the efficacy of RFA for the treatment of benign thyroid nodules, Chen et al. (2016) conducted a systematic review and meta-analysis and found that RFA was associated with a significant decrease in nodule volume at months 1, 3, 6, 12, and last follow-up. (38)

Fuller et al. (2014) reported on a systematic review of studies on RFA for benign thyroid tumors. (39) After RFA, statistically significant improvements were reported in combined symptom improvement and cosmetic scores on the 0 to 6 scale (mean, -2.96; 95% CI, -2.66 to -3.25) and withdrawal from methimazole (odds ratio, 40.34; 95% CI, 7.78 to 209.09). Twelve adverse events were reported, 2 of which were considered significant but did not require hospitalization.

Table 3 includes a comparison of studies included in the systematic reviews; the analyses by Cho et al. (2020) and Zu et al. (2024) contain the fewest number of included studies as a minimum follow-up duration of 3 years and 5 years, respectively were required for inclusion. Table 4 summarizes the characteristics of the systematic reviews and Table 5 contains the available results for nodule size reduction and complication rates. All of the systematic reviews are limited by high heterogeneity, inclusion of mostly single-center retrospective and/or noncontrolled studies, and generalizability concerns as included studies were mainly conducted in the Republic of Korea and Italy. They are further limited by a lack of comparison to surgical excision or PEI.

Table 3. Comparison of Meta-Analyses Assessing Radiofrequency Ablation for Benign Thyroid Nodules

Study	Xu (2024)* (36)	Cho (2020) * (37)	Chen (2016) (38)	Fuller (2014) (39)
Li (2022)	●			
Bernardi (2020)	●			
Aldea Martinez (2019)		●		

Deandrea (2019)	●	●		
Jung (2018)		●		
Sim (2017)		●		
Cesareo (2015)			●	
Sung (2015)			●	
Che (2015)			●	
Ugurlu (2015)			●	
Ji Hong (2015)			●	
Valcavi (2015)			●	
Bernardi (2014)			●	
Turtulici (2014)			●	
Yoon (2014)			●	
Lim (2013)	●		●	
Ha (2013)			●	
Sung (2013)			●	●
Huh (2012)				●
Faggiano (2012)			●	●
Jang (2012)				●
Kim (2012)				●
Baek (2010)			●	●
Lee (2010)			●	
Spiezia (2009)			●	●
Jeong (2008)			●	
Deandrea (2008)			●	●
Kim (2006)			●	●

* Only studies addressing radiofrequency ablation are included.

Table 4. Characteristics of Meta-Analyses Assessing Radiofrequency Ablation for Benign Thyroid Nodules

Study	Dates	Trials	Participants	N (Range)	Design	Duration
Xu (2024) (36)	Through Feb 2023	5	Patients with a benign thyroid nodule treated with thermal ablation (RFA [3 studies], microwave [1 study] or laser [2 studies])	939 (20 to 406)	5 retrospective cohorts	Approximately 5 years
Cho (2020) (37)	2010-2019	12	Patients with a benign thyroid nodule treated with thermal	1208 (24 to 276)	2 prospective and 10 retrospective cohorts	At least 3 years

			ablation (RFA [5 studies] or laser [7 studies])			
Chen (2016) (38)	2006-2016	20	Patients with a benign thyroid nodule treated with RFA	1090 (11 to 236)	Prospective and retrospective cohorts	Varied, 6 to 49.4 months
Fuller (2014) (39)	2006-2013	9	Patients with a benign thyroid nodule treated with RFA	284 (15 to 94)	Prospective studies (5 observational, 4 randomized trials)	Varied, 3 to 12 months

RFA: radiofrequency ablation.

Table 5. Key Results of Meta-Analyses Assessing Radiofrequency Ablation for Benign Thyroid Nodules

Study	Reduction in nodule size from baseline	Complication rate
Cho (2020) (37)	Relative volume reduction, VRR	
Total N, nodules (patients)	695 (680)	695 (680)
Pooled effect (95% CI)	6 mo: 64.5% (56.1% to 72.1%) 12 mo: 76.9% (65% to 85.7%) 24 mo: 80.1% (66.4% to 89.2%) 36 mo: 80.3% (66% to 89.5%)	4.6%
I^2 (p)	73.7%-95.9%	
Chen (2016) (38)	Absolute volume reduction, SMD	
Total N, nodules (patients)	1406 (1090)	
Pooled effect (95% CI)	1 mo: 0.83 (0.47 to 1.19) 3 mo: 1.31 (0.76 to 1.85) 6 mo: 1.25 (0.90 to 1.59) 12 mo: 4.16 (2.25 to 6.07)	
I^2 (p)	90.3%-98.7%	
Fuller (2014) (39)	Absolute volume reduction, SMD (follow up time frame not specified)	
Total N, nodules (patients)	284 (276)	
Pooled effect (95% CI)	-9.77 mL (-13.83 to -5.72)	
I^2 (p)	98% (<0.00001)	
Xu (2024)* (36)	Volume reduction rate	Regrowth
Total N, nodules (patients)	NR (939)	
Pooled effect (95% CI)	74.48% (70.05 to 78.91)	10.60% (1.50 to 19.80)

CI: confidence interval; NR: not reported; SMD: standard mean difference; VRR: volume reduction rate; mo: months.

*Includes data for all patients; RFA not separately analyzed from other methods of thermal ablation.

Section Summary: Benign Thyroid Tumors

Evidence on the treatment of benign thyroid nodules includes randomized and nonrandomized trials, case series, and systematic reviews of these studies. Systematic reviews have demonstrated that RFA results in a significant reduction in thyroid nodule size with a 2020 review showing that these changes remain durable through at least 36 months and a 2024 review indicating durability up to 5 years. Complication rates are generally low but include voice changes. The data are limited by significant heterogeneity in meta-analyses, a lack of generalizability to populations outside Republic of Korea and Italy, and a lack of comparators more relevant to practice in the United States.

Miscellaneous Solid Tumors

Clinical Context and Therapy Purpose

RFA has been investigated for use in individuals with different lesions in different anatomic sites. These anatomic sites include but are not limited to the thyroid, pancreas, and head and neck.

In individuals with head and neck cancer with recurrent disease, surgical salvage attempts are poor in terms of local control, survival, and QOL; further, these recurrent tumors are often untreatable with standard salvage therapies. Palliative chemotherapy or comfort measures may be offered. The safety and efficacy of RFA have been investigated as an option for palliative treatment in these situations.

The purpose of RFA in patients who have miscellaneous solid tumors is to provide a treatment option that is an alternative to or an improvement on existing therapies.

The following PICO was used to select literature to inform this policy.

Populations

The relevant population of interest is individuals with miscellaneous solid tumors (e.g., head and neck, thyroid cancer, pancreas).

Interventions

The therapy being considered is RFA.

Comparators

The following practices are currently being used to treat miscellaneous solid tumors: surgical excision or other local treatments specific to the tumor type.

Outcomes

The general outcomes of interest vary by disease state but include OS, tumor recurrence, and reductions in pain.

Patient follow-up will vary by disease state.

Study Selection Criteria

Methodologically credible studies were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.
- To assess long-term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Thyroid Cancer

Kim et al. (2015) reported on a comparative review of 73 patients with recurrent thyroid cancer smaller than 2 cm who had been treated with RFA (n=27) or repeat surgery (n=46). (40) RFA was performed in cases of patient refusal to undergo surgery or poor medical condition. Data were weighted to minimize potential confounders. The 3-year recurrence-free survival rates were similar for RFA (92.6%) and surgery (92.2%, p=.681). Posttreatment hoarseness rate did not differ between the RFA (7.3%) and surgery (9.0%) groups. Posttreatment hypocalcemia occurred only in the surgery group (11.6%).

Head and Neck Cancer

Owen et al. (2011) reported on RFA for 13 patients with recurrent and/or unresectable head and neck cancer who failed curative treatment. (41) Median patient survival was 127 days. While the stable disease was reported in 8 patients after RFA, and QOL scores improved, 3 deaths occurred (1 carotid hemorrhage, 2 strokes).

A case series of RFA for 14 patients with recurrent advanced head and neck malignancies was reported by Brook et al. (2008). (42) Tumor targeting and electrode deployment were successful in all cases, and 4 of 6 patients who completed QOL assessments showed improvement. Three major complications (in 27 [11%] applications) occurred 7 days to 2 weeks postprocedure. They included stroke, carotid artery rupture leading to death, and threatened carotid artery rupture with subsequent stroke. Retrospective analysis of intraprocedural CT scans revealed that the retractable electrodes were within 1 cm of the carotid artery during ablation in these cases.

A case series by Owen et al. (2004) showed that palliative CT-guided RFA provided subjective improvement with regard to pain, appearance, and function in 12 patients who had recurrent and advanced head and neck malignancies and were not candidates for radiotherapy or surgery. (43) The procedure appeared reasonably safe and feasible for this indication.

Uterine Myomas

A prospective observational study by Rey et al. (2019) assessed the effectiveness of transvaginal ultrasound-guided RFA of myomas in reducing tumor volume and eliminating menorrhagia associated with myomas. (44) The study included 205 women with symptomatic type II/III

uterine submucosal or intramural cavity-distorting myomas undergoing RFA. The preoperative mean (standard deviation) volume of the myomas was 122.4 (182.5) cm³ (95% CI, 82.1 to 162.8). Mean myoma volume decreased significantly at 1 (85.2 [147.9] cm³; p=.001), 3 (67.3 [138.0] cm³; p=.001), 6 (59.3 [135.3] cm³; p=.001), and 12 months (49.6 [121.4] cm³; p=.001). At 12 months, the mean volume reduction was 60% compared with preoperative volume. All patients returned to normal menstruation at a mean follow-up of 3 months and 12 months. Of the 205 patients, 201 (98.04%) were satisfied with the procedure. The investigators conceded that a larger population with a longer follow-up is needed but their study suggests that transvaginal ultrasound-guided RFA of myomas is effective and safe for treating select patients with metrorrhagia secondary to myomas.

In a large series, Yin et al. (2015) evaluated the effectiveness and safety of RFA for uterine myomas in a 10-year retrospective cohort study. (45) From 2001 to 2011, a total of 1216 patients treated for uterine myomas were divided into 2 groups. Group A consisted of 476 premenopausal patients (average age, 36 years) who had an average of 1.7 myomas with an average diameter of 4.5 cm. Group B consisted of 740 menopausal patients (average age, 48 years) with an average of 2.6 myomas with an average diameter of 5.0 cm. Patients were followed for a mean of 36 months. At 1, 3, 6, 12, and 24 months after RFA, the average diameters of myomas in group A were 3.8, 3.0, 2.7, 2.4, and 2.2 cm, respectively; 48% (227/476) of patients had a residual tumor at 12 months. In group B, myoma diameters were 4.7, 3.7, 3.3, 2.3, and 2.3 cm, respectively; 59% (435/740) of patients had trace disease at 12 months. Three months after RFA treatment, myoma volumes were significantly reduced in both groups (p<.01), although group B had a higher rate of residual tumor 12 months after RFA than group A (p<.05). Clinical symptoms and health-related QOL were significantly improved after RFA in both groups. The postoperative recurrence rate of uterine myomas was significantly higher in group A at 10.7% (51/476) than in group B at 2.4% (18/740; p<.05).

Adrenal Tumors

Liu et al. (2020) retrospectively evaluated the clinical outcomes of percutaneous ultrasound-guided RFA in the treatment of adrenal metastasis as compared to adrenalectomy. (46) Of the 60 patients included, 29 received RFA and 31 received adrenalectomy. The first technical success rate for RFA was 72.4%; 5 of the 8 patients had a repeat RFA and 4 of those achieved a complete response. In the adrenalectomy group, all patients achieved a R0 resection. Major complications were reported in 1 patient in the RFA group (ventricular fibrillation) and 2 patients in the adrenalectomy group (ascites, surgical site infection). The 1-, 2-, and 3-year local tumor progression rates after RFA were 17.1%, 30.9% and 44.7%, respectively, compared to 6.5%, 6.5% and 6.5% in adrenalectomy group (p=.028). There was no significant difference between groups for mean OS (2.3 ± 0.3 years for RFA and 3.9 ± 0.6 years for adrenalectomy, p=.057). Limitations of the study include its retrospective design, potential selection bias on which patients received each treatment, and a high prevalence of patients with adrenal metastasis secondary to hepatocellular carcinoma, which exceeded the expected number of cases based on global prevalence rates.

Liu et al. (2016) retrospectively compared laparoscopic adrenalectomy with CT-guided percutaneous RFA for the treatment of aldosterone-producing adenoma, evaluating short-term and long-term outcomes of normalized aldosterone-to-renin ratio, hypokalemia, and hypertension. (18) Of 63 patients, 27 were in the laparoscopic adrenalectomy group and 36 were in the RFA group. Primary aldosteronism was seen in 33 of 36 patients treated with RFA and all 27 who had laparoscopic adrenalectomy ($p=.180$), within a median follow-up of 5 to 7 years. RFA was associated with faster recovery postprocedure, but hypertension was less frequently resolved using RFA (13/36 patients) compared with laparoscopic adrenalectomy (19/27 patients; $p=.007$). The use of posture test and CT for subtype classification of primary aldosteronism is the major limitation of the study, as well as the retrospective design.

Retrospectively, Yang et al. (2016) compared the efficacy and safety of RFA with laparoscopic adrenalectomy in treating aldosterone-producing adenoma of the adrenal gland. (47) From 2009 to 2013, 25 patients diagnosed with unilateral adrenal aldosterone-producing adenoma and similar tumor size (<25 mm) were allocated to a control group ($n=18$) that underwent laparoscopic adrenalectomy and a test group ($n=7$) that underwent CT-guided percutaneous RFA. Complete tumor ablation on follow-up CT scan and normalization of serum aldosterone-to-renin were the primary outcomes compared in this study. Success in the RFA group reached 100% within 3 to 6 months, compared with 94.4% in the laparoscopic adrenalectomy group, and normalization ability was statistically equivalent in both groups. The study's retrospective design and small sample are the main limitations of this study.

Other Tumors

A single-arm, retrospective, paired-comparison study by Locklin et al. (2004) evaluated the short-term efficacy of RFA in reducing pain and improving function in patients with unresectable, painful soft tissue neoplasms recalcitrant to conventional therapies. (48) Patients had tumors located in a variety of sites including chest wall, pelvis, breast, perirectal, renal, aortocaval, retroperitoneal, and superficial soft tissues. All had failed conventional methods of palliation or experienced dose-limiting adverse events from pain medication. Although not all Brief Pain Inventory scores were statistically significant, all mean scores trended down over time after ablation. Complications from RFA were minor or insignificant in all but 1 patient who had skin breakdown and infection of an ablated superficial tumor site.

Additional research has addressed the use of RFA in solid malignancies (49, 50) and in the pancreas. (51-53) A systematic review by Rombouts et al. (2015) has examined studies of ablative therapies, including RFA, in patients with locally advanced pancreatic cancer. (54) No RCTs were identified, and conclusions were limited by the sparse evidence available on RFA in this setting.

Stereotactic radiofrequency thermocoagulation for epileptogenic hypothalamic hamartomas was described in a retrospective analysis by Kameyama et al. (2009) who evaluated 25 patients with gelastic seizures (a rare type of seizure). (55) Other seizure types were exhibited in 22 (88.0%) patients, precocious puberty in 8 (32.0%), behavioral disorder in 10 (40.0%), and mental disability in 14 (56.0%). Gelastic seizures resolved in all but 2 patients. Complete seizure

freedom was achieved in 19 (76.0%) patients. These patients experienced resolution of all seizure types and behavioral disorder and also demonstrated intellectual improvement.

Preliminary results of endoscopic RFA of rectosigmoid tumors have been described by Vavra et al. (2009). (56) Twelve patients were treated with the Endoblate RFA device, with 10 patients having surgical resection after ablation. Histology of the resected specimens showed that, on average, 82% (range, 60% to 99%) of the tumor mass was destroyed in the ablation zone.

Small case series on RFA for colorectal and rectal carcinoma have demonstrated a debulking role for RFA. (57, 58) These case series did not permit comparison with an available alternative.

Section Summary: Miscellaneous Solid Tumor

Evidence on the use of RFA to treat other types of solid tumors consists of a small number of case series, prospective studies, or retrospective comparative studies. Reporting on outcomes is limited. The evidence base does not support a conclusion on the effects of RFA for the tumor types included in this evidence review.

Summary of Evidence

For individuals who have painful osteolytic bone metastases who have failed or are poor candidates for standard treatments who receive radiofrequency ablation (RFA), the evidence includes a prospective cohort study and case series. Relevant outcomes are symptoms, change in disease status, quality of life (QOL), medication use, and treatment-related morbidity. A prospective cohort study and case series have shown clinically significant pain relief (defined as a decrease of 2 units from baseline on the Brief Pain Inventory scale) or reduction in opioid use following treatment of painful osteolytic metastases. A multicenter, prospective study reported significant reductions in pain through the 6-month follow-up period, with 59% of patients achieving immediate improvement in pain within 3 days of RFA. The population is comprised of patients with few or no treatment options, for whom short-term pain relief is an appropriate clinical outcome. The evidence is sufficient to determine that the technology results in an improvement in the net health outcome.

For individuals who have painful osteoid osteomas who receive RFA, the evidence includes numerous observational studies and systematic reviews of these studies. Relevant outcomes are symptoms, change in disease status, QOL, medication use, and treatment-related morbidity. In a systematic review of thermal ablation techniques, clinical success (pain-free) was achieved in 94% to 98% of patients. Most patients (89% to 96%) remained pain-free when assessed during longer-term follow-up. Another systematic review reported similar success rates noting an average 8.3% failure rate among patients receiving computed tomography (CT)-guided RFA. Although no randomized trials of RFA for osteoid osteomas have been performed, uncontrolled studies have demonstrated RFA can provide adequate symptom relief with minimal complications for a population for whom short-term symptom relief and avoidance of invasive procedures are appropriate clinical outcomes. The evidence is sufficient to determine that the technology results in an improvement in the net health outcome.

For individuals who have localized renal cell carcinoma (RCC) that is no more than 4 cm in size who receive RFA, the evidence includes a randomized controlled trial (RCT), numerous observational studies, and systematic reviews of these studies. Relevant outcomes are overall survival (OS), change in disease status, QOL, and treatment-related morbidity. A recent meta-analysis that included only an RCT and cohort studies found that RFA was as effective as nephrectomy for small renal tumors, with a reduction in complications. Another recent meta-analysis found that partial nephrectomy (PN) was superior to ablative techniques (the study included RFA but also cryoablation and microwave ablation) in overall mortality and local recurrence but not in cancer-specific mortality. It also found fewer complications and improved renal function with ablation. A meta-analysis from 2022 found that PN was superior to ablation (RFA, cryoablation, and microwave ablation) in local recurrence. Overall complications, decline in renal function, and cancer-specific mortality rates did not differ between ablation and PN. A 2025 meta-analysis found that RFA achieved high local control and cancer-specific survival rates comparable to other ablative therapies, although SBRT showed superior local control for larger tumors. Although inconsistent, the evidence does suggest that, for small renal tumors, RFA may result in a similar rate of disease progression with a lower complication rate than nephrectomy. However, comparative trials are needed to determine with greater certainty the effects of these treatments in the same patient population. The evidence is insufficient to determine that the technology results in an improvement in the net health outcome.

For individuals who have inoperable primary pulmonary tumors or nonpulmonary tumors metastatic to the lung who receive RFA, the evidence includes prospective observational studies and systematic reviews of these studies. Relevant outcomes are OS, change in disease status, QOL, and treatment-related morbidity. A multicenter study found that for tumors less than 3.5 cm in size, RFA can lead to a complete response in as many as 88% of patients for at least 1 year. Two-year survival rates have been reported to range from 41% to 75% in case series, with 5-year survival rates of 20% to 27%. In general, the evidence suggests that RFA results in adequate survival and tumor control in patients who are not surgical candidates, with low morbidity rates. The evidence is sufficient to determine that the technology results in an improvement in the net health outcome.

For individuals who have breast tumors who receive RFA, the evidence includes observational studies and systematic reviews of these studies. Relevant outcomes are OS, change in disease status, QOL, and treatment-related morbidity. Evidence has reported varied and incomplete ablation rates with concerns about postablation tumor cell viability. Long-term improvements in health outcomes have not been demonstrated. Additionally, available studies do not permit comparisons with conventional breast-conserving procedures. Further prospective studies, with long-term follow-up, should focus on whether RFA of the breast for small tumors can provide local control and survival rates compared with conventional breast-conserving treatment. The evidence is insufficient to determine that the technology results in an improvement in the net health outcome.

For individuals who have benign thyroid tumors who receive RFA, the evidence includes RCTs, prospective studies, case series, and systematic reviews of these studies. Relevant outcomes

are symptoms, change in disease status, QOL, medication use, and treatment-related morbidity. Systematic reviews have demonstrated that RFA results in a significant reduction in thyroid nodule size, with a 2020 review showing that these changes remain durable through at least 36 months and a 2024 review indicating durability up to 5 years. Complication rates are generally low but include voice changes. The data are limited by significant heterogeneity in meta-analyses, a lack of generalizability to populations outside Republic of Korea and Italy, and a lack of comparators more relevant to practice in the United States. Further studies comparing RFA to percutaneous ethanol injection (PEI) or surgery would be more informative in determining the potential utility of RFA in patients with symptomatic or large benign thyroid tumors as these are the recommended treatment options per the American Thyroid Association. The evidence is insufficient to determine that the technology results in an improvement in the net health outcome.

For individuals who have miscellaneous tumors (e.g., head and neck, thyroid cancer, pancreas) who receive RFA, the evidence includes a few case series, prospective observational studies, and retrospective comparative studies. Relevant outcomes are OS, change in disease status, QOL, and treatment-related morbidity. There is a limited evidence base for these tumor types. Reporting on outcomes or comparisons with other treatments is limited. These studies do not permit conclusions on the health benefits of RFA. The evidence is insufficient to determine that the technology results in an improvement in the net health outcome.

Practice Guidelines and Position Statements

American College of Chest Physicians

The American College of Chest Physicians (2025) guidelines on management of patients with early-stage non-small cell lung cancer state that "image-guided thermal ablation therapies such as radiofrequency ablation (RFA), microwave ablation (MWA), and cryoablation (CA) have also been used to treat lung tumors, particularly in patients with contraindications to surgery. For patients deemed to have "high operative risk" (i.e., those who cannot tolerate lobectomy, but are candidates for sublobar resection) stage I NSCLC, the appropriateness of potential alternatives [e.g., SBRT; ablation] to surgery is an active area of investigation." (59) They also state that "To date, no specific technique of ablation (radiofrequency, microwave, or cryoablation) has proven superior, and therefore outcomes from various modalities are most often combined to evaluate local control and overall survival."

American Head and Neck Society - Endocrine Surgery Section

An international, multidisciplinary consensus statement on RFA and related ultrasound-guided ablation technologies for the treatment of benign and malignant thyroid disease was released in 2022 through a collaboration of international professional societies, including the Endocrine Surgery Section of the American Head and Neck Society. (60) Select relevant recommendations from the guideline are listed in Table 6.

Table 6. Summary of RFA Recommendations for Treatment of Benign and Malignant Thyroid Disease*

Recommendation 1	US-guided ablation procedures may be used as a first-line alternative to surgery for patients with benign thyroid nodules contributing to compressive and/or cosmetic symptoms.
Recommendation 2	Although less efficacious than surgery or RAI in normalizing thyroid function, thermal ablation procedures can be a safe therapeutic alternative in patients with an autonomously functional thyroid nodule and contraindications to first-line techniques.
Recommendation 3a	US-guided ablation procedures may be considered in patients with suitable primary papillary microcarcinoma who are unfit for surgery or decline surgery or active surveillance
Recommendation 3b	US-guided ablation procedures may be considered in patients with suitable recurrent papillary thyroid carcinoma who are unfit for surgery or decline surgery or active surveillance
Recommendation 3c	Repeat ablation of a benign nodule can be considered for remnant nodular tissue contributing to unresolved symptomatic or cosmetic concerns

*This is not a comprehensive list of recommendations from the guideline.

RAI: radioactive iodine; RFA: radiofrequency ablation; US: Ultrasound.

American Urological Association

The American Urological Association (AUA; 2017) guideline on renal masses and localized renal cancer affirms that partial nephrectomy should be prioritized for the management of cT1a renal masses when intervention is indicated. (61) Thermal ablation should be considered "as an alternate approach for the management of cT1a renal masses <3 cm in size." The guidelines were updated in 2021, and recommendations are generally consistent with what was published in the 2017 guideline. (62) The 2021 AUA guideline explicitly states that RFA and cryoablation may be offered as options to patients who elect thermal ablation.

American Thyroid Association

The American Thyroid Association (2015) guideline on the management of thyroid nodules and differentiated thyroid cancer provides recommendations for management. (35) Patients with a benign cytology diagnosis or those very unlikely to be malignant (e.g., purely cystic nodule) should undergo surveillance with the frequency determined by the level of suspicion for a missed malignancy. Medical or surgical intervention is considered if the nodules are large (>4 cm), causing compressive or structural symptoms, or if there is clinical concern. Recurrent cystic thyroid nodules with benign cytology should be considered for surgical removal or percutaneous ethanol injection. For differentiated thyroid cancer, "localized treatments with thermal (radiofrequency or cryo-) ablation, ethanol ablation, or chemoembolization may be beneficial in patients with a single or a few metastases and in those with metastases at high risk of local complications."

National Comprehensive Cancer Network

The National Comprehensive Cancer Network (NCCN) guidelines for the treatment of NSCLC (v.8.2025) state (63): "For medically operable disease, resection is the preferred local treatment

modality (other modalities include SABR [stereotactic ablative radiotherapy], thermal ablation such as radiofrequency ablation, and cryotherapy)." For patients who are not amenable to surgery, image-guided thermal ablation therapy (IGTA; includes RFA, microwave ablation, and cryoablation) may be considered. The guidance states "IGTA is an option for the management of NSCLC lesions <3 cm. Ablation for NSCLC lesions >3 cm may be associated with higher rates of local recurrence and complications."

The NCCN guidelines for thyroid carcinoma (v.1.2025) indicate that local therapies such as RFA may be considered for locoregional recurrence of thyroid carcinoma-papillary carcinoma in select patients with limited burden nodal disease. Additionally, local therapies, including RFA, can be considered in those with metastatic disease. (64)

The NCCN guidelines (v.1.2026) for renal cell carcinoma indicate that "percutaneous ablation (e.g., cryosurgery, radiofrequency ablation, microwave ablation) is an option for the management of clinical stage T1 renal lesions. Percutaneous ablation is an option for clinical T1b masses in select patients not eligible for surgery. Biopsy of lesions is recommended to be done prior to or at time of ablation. Percutaneous ablation may require multiple treatments to achieve the same oncologic outcomes as conventional surgery." (65)

The NCCN colon cancer guidelines (v.4.2025) state that "resection is the standard approach for the local treatment of resectable metastatic disease. However, patients with liver or lung oligometastases can also be considered for tumor ablation therapy, particularly in cases that may not be optimal for resection." (66) "There is extensive evidence on the use of RFA as a reasonable treatment option for non-surgical candidates and for recurrent disease after hepatectomy with small liver metastases that can be treated with clear margins."

The NCCN guidelines for head and neck cancers (v.5.2025), (67) breast cancer (v.5.2025), (68) bone cancer (v.1.2026), (69) and pancreatic adenocarcinoma (v.2.2025) (70) do not mention RFA.

National Institute for Health and Care Excellence (NICE)

The NICE guidance (2004) on osteoid osteoma indicated that "current evidence on the safety and efficacy of computed tomography (CT)-guided thermocoagulation of osteoid osteoma appears adequate to support its use...." (71)

Updated NICE guidance (2010) on renal cancer has indicated that "evidence on the safety and efficacy of percutaneous radiofrequency ablation (RFA) ... in the short and medium term appears adequate to support the use of this procedure provided that patients are followed up in the long term." (70)

The NICE guidance (2010) on RFA for primary and secondary lung cancers has stated: "Current evidence on the efficacy of percutaneous radiofrequency ablation (RFA) ... is adequate in terms of tumor control." (72) The NICE also indicated RFA might "be used in patients with small, early-stage lung cancers or small numbers of lung metastases who are unsuitable for, or prefer not to

undergo, surgery. It may also have a place in multi-modality treatment of more advanced primary lung cancers." The guidance warned of serious complications (e.g., pneumothorax) among lung cancer patients.

The NICE guidance (2016) on benign thyroid nodules stated: "Current evidence on the safety and efficacy of ultrasound-guided percutaneous radiofrequency ablation ... is adequate to support the use of this procedure...." (73)

Society of Interventional Radiology

The Society of Interventional Radiology (2020) published a position statement on the role of percutaneous ablation in renal cell carcinoma. (74) Their relevant recommendations are as follows:

- "In patients with small renal tumors (stage T1a), percutaneous thermal ablation is a safe and effective treatment with fewer complications than nephrectomy and acceptable long-term oncological and survival outcomes. (Level of Evidence: C; Strength of Recommendation: Moderate)"
- "In selected patients with suspected T1a renal cell carcinoma, percutaneous thermal ablation should be offered over active surveillance. (Level of Evidence: C; Strength of Recommendation: Moderate)"
- "In high-risk patients with T1b renal cell carcinoma who are not surgical candidates, percutaneous thermal ablation may be an appropriate treatment option; however, further research in this area is required. (Level of Evidence: D; Strength of Recommendation: Weak)"
- "Radiofrequency ablation, cryoablation, and microwave ablation are all appropriate modalities for thermal ablation, and method of ablation should be left to the discretion of the operating physician. (Level of Evidence: D; Strength of Recommendation: Weak)"

Ongoing and Unpublished Clinical Trials

Some currently ongoing and/or unpublished trials that might influence this review are listed in Table 7.

Table 7. Summary of Key Trials

NCT Number	Trial Name	Planned Enrollment	Completion Date
NCT05189821	RFA Treatment for Papillary Thyroid Microcarcinoma Cohort	50	Nov 2027
NCT05189808	Radiofrequency Ablation for Indeterminate Bethesda III Thyroid Nodules	50	Aug 2027
NCT04619472	A Multicenter, Single Group Target Value Clinical Study to Evaluate Safety and Effectiveness of Radiofrequency Ablation System in the Treatment of Peripheral Lung Tumors	126	Mar 2023

NCT01051037	Phase II Study Evaluating Safety and Efficacy of Stereotactic Body Radiotherapy and Radiofrequency Ablation for Medically Inoperable and Recurrent Lung Tumors Near Central Airways	17	Dec 2017 (completed)
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NCT: national clinical trial.

Coding

Procedure codes on Medical Policy documents are included **only** as a general reference tool for each policy. **They may not be all-inclusive.**

The presence or absence of procedure, service, supply, or device codes in a Medical Policy document has no relevance for determination of benefit coverage for members or reimbursement for providers. **Only the written coverage position in a Medical Policy should be used for such determinations.**

Benefit coverage determinations based on written Medical Policy coverage positions must include review of the member's benefit contract or Summary Plan Description (SPD) for defined coverage vs. non-coverage, benefit exclusions, and benefit limitations such as dollar or duration caps.

CPT Codes	20982, 32998, 41530, 50541, 50542, 50592, 60660, 60661, 60699, 76940, 77013, 77022
HCPCS Codes	None

*Current Procedural Terminology (CPT®) ©2024 American Medical Association: Chicago, IL.

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Centers for Medicare and Medicaid Services (CMS)

The information contained in this section is for informational purposes only. HCSC makes no representation as to the accuracy of this information. It is not to be used for claims adjudication for HCSC Plans.

The Centers for Medicare and Medicaid Services (CMS) does not have a national Medicare coverage position. Coverage may be subject to local carrier discretion.

A national coverage position for Medicare may have been developed since this medical policy document was written. See Medicare's National Coverage at <<https://www.cms.hhs.gov>>.

Policy History/Revision

Date	Description of Change
12/01/2025	Document updated. The following change was made to Coverage: Moved information in NOTE 1 to Policy Guidelines section. Added references 13 and 59; others updated and some removed. Title changed from: Radiofrequency Ablation (RFA) of Solid Tumors, Excluding Liver.
01/01/2025	Document updated with literature review. Coverage unchanged. Reference 50 added; others updated.
12/01/2023	Reviewed. No changes.
04/15/2023	Document updated with literature review. The following changes were made to Coverage: 1) Renal Cell Carcinoma, RFA may be considered medically necessary to treat localized RCC that is no more than 4 cm in size when criteria 1 and 2 are met and 2) Non-Small-Cell Lung Cancer (NSCLC) Primary Pulmonary Tumor, RFA may be considered medically necessary to treat an isolated peripheral NSCLC (primary pulmonary) lesion that is no more than 3 cm in size when criteria 1 and 2 are met and 3) Nonpulmonary Tumor(s) Metastatic to the Lung, RFA may be considered medically necessary to treat malignant nonpulmonary tumor(s) metastatic to the lung that are no more than 3 cm in size when criteria 1 and 2 are met. References 1-5, 17, 38, 74, 76, 82, 83, 85 and 90 added.
02/01/2022	Reviewed. No changes.
10/15/2021	Document updated with literature review. Clarification to coverage with no change to intent. References 4, 9, 16, 17, 41, 42, 51, 53, 67 and 73 added; numerous references removed; others removed.
01/15/2021	Reviewed. No changes.
09/15/2020	Document updated with literature review. The following change was made to Coverage: Added “palliate pain in patients with” as a clarification to the statement on osteolytic bone metastases. References 5, 6, 10, 13-17, 28, 31-32, 37, 39, 47-48, 68 and 72 added; numerous references removed.
09/15/2019	Reviewed. No changes.
07/01/2018	Document updated with literature review. Osteoid Osteomas or Bony Metastases split into 2 separate coverage statements: 1) Radiofrequency ablation (RFA) may be considered medically necessary to treat osteolytic bone metastases in patients who have failed or are a poor candidate for standard treatment such as surgical intervention, radiation, chemotherapy, or opioids; and 2) RFA may be considered medically necessary to treat osteoid osteomas that cannot be managed successfully with medical treatment. Numerous wording changes to coverage, but intent of coverage remained unchanged. The following NOTE was added: NOTE 2: See medical policy SUR709.029 for Radiofrequency Ablation (RFA) of Primary or Metastatic Liver Tumors. The following references have been added: 8, 13, 34, and 63; numerous references removed.

04/15/2017	Reviewed. No changes.
08/01/2016	Policy updated with literature review. The following changes were made to Coverage: 1) Removed “only when determined to be indicated by the treating physician” from the osteoid osteoma or bony metastases coverage criteria; 2) Removed “palliative treatment of medically or surgically inoperable tumor(s)” and added “that is no more than 4 cm in size” to the renal cell carcinoma coverage criteria; 3) Revised bullet from “patients who have failed, or are a poor candidate for standard treatment such as surgical intervention, radiation, chemotherapy and opioids” to now state “the patient is not considered a surgical candidate” in the renal cell carcinoma coverage criteria; 4) Removed “radiofrequency ablation may be considered medically necessary for treatment of primary and metastatic neoplasms when removal of the neoplasm may be curative but the patient has been determined by the treating physician to be unable to tolerate surgical resection.” 5) Removed “Primary and metastatic neoplasms when the patient has been determined by the treating physician to be able to tolerate surgical resection” from the EIU coverage section.
05/15/2015	Reviewed. No changes.
07/01/2014	Document updated with literature review. Coverage of osteomas, bony metastases, and inoperable tumors is unchanged. In addition, the following changes were made: 1) Coverage of renal cell carcinoma (RCC) was moved to this policy from SUR710.017 (Radiofrequency Ablation [RFA] and Cryoablation of Renal Cell Carcinoma [RCC]) and SUR710.017 was deleted; coverage of RCC is unchanged. 2) Coverage of pulmonary tumors was moved from SUR706.012 (Radiofrequency Ablation [RFA] of Pulmonary Tumors) and SUR706.012 was deleted; 3) New detailed coverage criteria were added for isolated peripheral non-small-cell lung cancer and for malignant nonpulmonary tumors metastatic to the lung.
06/01/2013	Document updated with literature review. Coverage unchanged. Rationale was revised.
07/01/2011	Document updated with literature review. Coverage unchanged.
08/15/2009	Revised/updated entire document with literature search. No change in coverage position.
06/01/2007	Revised/updated entire document
01/01/2007	New medical document

Appendix

Table A1. Comparison of Studies Included in SRs & M-As Assessing Radiofrequency Ablation for Renal Masses

Study	El Dib (2012) ^a (17)	Katsanos (2014) (16)	Uhlig (2019) (15)	Yanagisawa (2022) (14)
Pavlovich (2002)	●			

Gervais (2003)	●			
Lewin (2004)	●			
DiMarco (2004)	●			
Desai (2005)				●
Memarsadeghi (2006)	●			
Hegarty (2006)			●	
Zagoria (2007)	●			
Klinger (2007)	●			
Breen (2007)	●			
Stern (2007)		●	●	●
Bensalah (2007)			●	
Omalley (2007)			●	
Wingo (2008)	●			
Watane (2008)	●			
Raman (2008)	●			
Foyil (2008)			●	
Lucas (2008)			●	
Ko (2008)				●
Bird (2009)		●	●	
Lughezzani (2009)			●	
Turna (2009)			●	
Raman (2010)			●	
Takaki (2010)		●	●	
Klatte (2011)			●	●
Haber (2011)			●	
Sung (2012)		●	●	
Guan (2012)		●	●	●
Olweny (2012)		●	●	●
Haramis (2012)			●	●
Guillotreau (2012)			●	●
Whitson (2012)				●
Arnoux (2013)			●	
Atwell (2013)			●	
Panumatrassamee (2013)			●	
Psutka (2013)			●	
Tanagho (2013)			●	
Youn (2013)			●	
Emara (2014)			●	
Chang, Liu (2015)			●	●

Chang, Zhang (2015)			●	●
Cooper (2015)			●	
Danzig (2015)			●	
Kim (2015)			●	●
Desai (2015)			●	
Thompson (2015)			●	
Weinberg (2015)			●	
Fosatti (2015)				●
Ji (2016)			●	
Hinshaw (2016)			●	
Chehab (2016)			●	
Camacho (2016)			●	
Pantelidou (2016)			●	
Zechlinski (2016)			●	
Evans (2017)			●	
Caputo (2017)			●	●
Abboud (2017)			●	
Alekseev (2017)			●	
Bhindi (2017)			●	
Mason (2017)			●	
Liu (2017)			●	●
Wan (2017)			●	
Park (2018)				●
Park (2019)				●
Yanagisawa (2018)				●
Anglickis (2019)				●
Andrews (2019)				●
Acosta Ruiz (2019)				●
Rembeyo (2019)				●
Shapiro (2019)				●
Yu (2020)				●
Lesnyak (2020)				●
Garcia (2021)				●
Liu (2021)				●