Microwave ablation (MWA) of primary and metastatic tumors is considered investigative.

**Related Policies**

- **7.01.95**  
  Radiofrequency Ablation of Miscellaneous Solid Tumors Excluding Liver Tumors

- **7.01.526**  
  Cryosurgical Ablation of Miscellaneous Solid Tumors Other Than Liver, Prostate, or Dermatologic Tumors

- **8.01.505**  
  Transcatheter Arterial Chemoembolization as a Treatment for Primary or Metastatic Liver Malignancies

- **8.01.521**  
  Radioembolization for Primary and Metastatic Tumors of the Liver

**Policy Guidelines**

There are no CPT codes specific to microwave ablation.

According to a 2012 American Medical Association publication (*Clinical Examples in Radiology*, Vol. 8, Issue 3; Summer 2012), “microwave is part of the radiofrequency spectrum, and simply uses a different part of the radiofrequency spectrum to develop heat energy to destroy abnormal tissue.” Therefore, AMA recommends that microwave ablation should be reported using the CPT codes for radiofrequency ablation as noted in the coding table below.

When there is no specific CPT code for ablation, the unlisted CPT code for the anatomic area should be reported.
Microwave ablation (MWA) also referred to as microwave coagulation therapy destroys tumors and soft tissue using microwave energy. MWA is an option used to treat tumors not appropriate for resection or to treat patients ineligible for surgery due to age, comorbidities, or poor general health. MWA may be performed under sedation or anesthesia as an open procedure, laparoscopically, percutaneously, or thoracoscopically under image guidance.

Background
MWA is a technique in which the use of microwave energy induces an ultra-high speed, 915 MHz or 2.450 MHz (2.45 GHz), alternating electric field, which causes water molecule rotation and the creation of heat. This results in thermal coagulation and localized tissue necrosis. In MWA, a single microwave antenna or multiple antennas connected to a generator are inserted directly into the tumor or tissue to be ablated; energy from the antennas generates friction and heat. The local heat coagulates the tissue adjacent to the probe, resulting in a small, approximately 2- to 3-cm elliptical area (5 x 3 cm) of tissue ablation. In tumors greater than 2 cm in diameter, 2 to 3 antennas may be used simultaneously to increase the targeted area of MWA and shorten operative time. Multiple antennas may also be used simultaneously to ablate multiple tumors. Tissue ablation occurs quickly, within 1 minute after a pulse of energy, and multiple pulses may be delivered within a treatment session, depending on the size of the tumor. The cells killed by MWA are typically not removed but are gradually replaced by fibrosis and scar tissue. If there is local recurrence, it occurs at the edges. Treatment may be repeated as needed. MWA may be used to:
1. Control local tumor growth and prevent recurrence
2. Palliate symptoms
3. Extend survival duration

MWA is similar to radiofrequency (RFA) and cryosurgical ablation. However, MWA has potential advantages over RFA and cryosurgical ablation. In MWA, the heating process is active, which produces higher temperatures than the passive heating of RFA and should allow for more complete thermal ablation in less time. The higher temperatures reached with MWA (>100°C) can overcome the “heat sink” effect in which tissue cooling occurs from nearby blood flow in large vessels, potentially resulting in incomplete tumor ablation. MWA does not rely on the conduction of electricity for heating and, therefore, does not flow electrical current through patients and does not require grounding pads, because there is no risk of skin burns. Additionally, MWA does not produce electric noise, which allows ultrasound guidance during the procedure without interference, unlike RFA. Finally, MWA can take less time than RFA, because multiple antennas can be used simultaneously.

Complications from MWA are usually considered mild and may include pain and fever. Other potential complications associated with MWA include those caused by heat damage to normal tissue adjacent to the tumor (e.g., intestinal damage during MWA of the kidney or liver), structural damage along the probe track (e.g., pneumothorax as a consequence of procedures on the lung), liver enzyme elevation, liver abscess, ascites, pleural effusion, diaphragm injury or secondary tumors if cells seed during probe removal. MWA should be avoided in pregnant patients because potential risks to the patient and/or fetus have not been established and in patients with implanted electronic devices such as implantable pacemakers that may be adversely affected by...
microwave power output.

MWA was first used percutaneously in 1986 as an adjunct to liver biopsy. Since then, MWA has been used to ablate tumors and tissue to treat many conditions including: hepatocellular carcinoma, breast cancer, colorectal cancer metastatic to the liver, renal cell carcinoma, renal hamartoma, adrenal malignant carcinoma, non-small-cell lung cancer, intrahepatic primary cholangiocarcinoma, secondary splenomegaly and hypersplenism, abdominal tumors, and other tumors not amenable to resection. Well-established local or systemic treatment alternatives are available for each of these malignancies. The potential advantages of MWA for these cancers include improved local control and those common to any minimally invasive procedure (eg, preserving normal organ tissue, decreasing morbidity, shortening length of hospitalization). MWA also has been investigated as a treatment for unresectable hepatic tumors, both as primary treatment, palliative treatment, and as a bridge to liver transplant. In the latter setting, MWA is being assessed to determine whether it can reduce the incidence of tumor progression while awaiting transplantation and thus maintain a patient’s candidacy while awaiting liver transplant.

Regulatory Status

There are several devices cleared for marketing by FDA through the 510(k) process for MWA. Covidien’s (a subsidiary of Tyco Healthcare) Evident™ Microwave Ablation System has 510(k) clearance for soft tissue ablation, including partial or complete ablation of non-resectable liver tumors. The following devices have 510(k) clearance for MWA of (unspecified) soft tissue:

- BSD Medical Corporation’s MicroThermX® Microwave Ablation System (MTX-180);
- Microsulis Medical’s (now part of AngioDynamics) Acculis® Accu2i; and
- MicroSurgeon Microwave Soft Tissue Ablation Device;
- NeuWave Medical’s Certus 140™
- Valleylab’s (a subsidiary of Covidien) VivaWave® Microwave Ablation System;
- Vivant’s (acquired by Valleylab in 2005) Tri-Loop™ Microwave Ablation Probe;

These devices are considered substantially equivalent to previously FDA-approved radiofrequency and MWA devices. FDA product code: NEY.

This policy does not address MWA for the treatment of splenomegaly or ulcers or as a surgical coagulation tool.

Scope

Medical policies are systematically developed guidelines that serve as a resource for Company staff when determining coverage for specific medical procedures, drugs or devices. Coverage for medical services is subject to the limits and conditions of the member benefit plan. Members and their providers should consult the member benefit booklet or contact a customer service representative to determine whether there are any benefit limitations applicable to this service or supply. This medical policy does not apply to Medicare Advantage.

Benefit Application

N/A

Rationale

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### Breast Cancer
A 2010 review of ablation techniques by Zhao et al. for breast cancer found only 0% to 8% of breast tumors were completely ablated with microwave ablation (MWA). (1) The authors noted the studies identified for the review were mostly feasibility and pilot studies conducted in research settings.

In 2012, W. Zhou et al. reported on 41 patients treated with MWA directly followed by mastectomy for single breast tumors with a mean volume of 5.26±3.8 cm (range, 0.09-14.14 cm). (2) Complete tumor ablation was found by microscopic evaluation in 37 of the 41 tumors ablated (90%; 95% confidence interval [CI], 76.9% to 97.3%). Reversible thermal injuries to the skin and pectoralis major muscle occurred in 3 patients.

### Hepatocellular Carcinoma

#### Systematic Reviews
Chinnaratha et al published a meta-analysis of randomized controlled trials (RCTs) and observational studies that compared the effectiveness and safety of radiofrequency ablation (RFA) to MWA in patients with primary hepatocellular carcinoma (HCC). (3) MEDLINE, EMBASE, and Cochrane Central databases were searched between January 1980 and May 2014 for human studies comparing the 2 technologies. The primary outcome was the risk of local tumor progression (LTP); secondary outcomes were complete ablation, overall survival (OS), and major adverse events. Odds ratios (ORs) were combined across studies using a random-effects model. Ten studies (2 prospective, 8 retrospective) were included. The overall LTP rate was 14% (176/1298). There was no difference in LTP rates between RFA and MWA (OR=1.01; 95% CI, 0.67 to 1.50; p=0.9). The complete ablation rate, 1- and 3-year OS, and major adverse events were similar between the 2 modalities (p>0.05 for all).

Subgroup analysis showed LTP rates were lower with MWA for treatment of larger tumors (OR=1.83; 95% CI, 1.10 to 3.23; p=0.02). No significant publication bias was detected nor was interstudy heterogeneity ($I^2$<50%, p>0.1) observed for any measured outcomes.

Bertot et al conducted a systematic review in 2011 of ablation techniques for primary and secondary liver tumors. (4) This review included 2 studies (5,6) using MWA (total N=1185 patients). Pooled analysis was performed using a random effects model because of significant study heterogeneity. The pooled mortality rate for MWA was 0.23% (95% CI, 0.0% to 0.58%). The pooled rate for major complications following MWA was 4.6%.

In 2009, Ong et al. conducted a systematic review of studies on MWA for primary and secondary liver tumors. (7) Based on the results from 25 clinical studies reporting outcomes on MWA, the authors concluded MWA is an effective and safe technique for liver tumor ablation with low complication rates and survival rates comparable with hepatic resection. However, rates of local recurrence after MWA were noted to be higher than hepatic resection. In most studies, HCC recurrence rates were approximately 10% but were also noted to be as high as 50%, which the authors indicated can be addressed with further ablation. Survival rates in the studies on MWA for HCC were as high as 92% at 3 years and 72% at 5 years, which was noted to be comparable with RFA and percutaneous ethanol injections. Pain and fever were the most frequently reported complications, but complications increased when there were more tumors, larger tumors, and more microwave antennas used.

#### Comparative Studies
No RCTs comparing MWA to RFA were identified. The available studies are nonrandomized comparisons, and all except 1 study is retrospective.

Abdelaziz et al reported a prospective study in 2015 that evaluated the efficacy and safety of MWA and transarterial chemoembolization (TACE) for large tumors (5-7 cm) and assessed their effects on local tumor progression and survival. (8) Sixty-four patients with large lesions were divided into 2 groups treated by MWA or by TACE. Both groups were comparable in demographic and ultrasonographic tumor features. MWA completely

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The evidence review was created in 2011 and has been periodically updated with literature searches of the MEDLINE database. The last literature review was conducted through February 15, 2016. The findings of the literature review are summarized next with select studies.
ablated 75% of cases in fewer sessions, with a lower incidence of tumor recurrence (p=0.02), development of de novo lesions (p=0.03), occurrence of posttreatment ascites (p=0.003), and had higher OS rates (p=0.04) than TACE. Mean OS in the MWA group was 22 months and 14 months in the TACE group. Actuarial probabilities of survival at 12 and 18 months were 78% and 68%, respectively, in the MWA group and 52% and 29%, respectively, in the TACE group.

Vogl et al published a retrospective comparative study in 2015. (9) It enrolled 53 patients with 68 liver lesions due to HCC. MWA was performed in 36 patients and RFA in 32 patients. There were no differences between groups on complete response immediately following treatment or for progression-free survival at 12 month or OS at 3 years. In 2013, Ding et al a retrospectively compared 113 patients treated with MWA for 131 HCC tumors and 85 patients treated with RFA for 98 HCC tumors. (10) Rates of complete ablation, local recurrence, disease-free (DFS) and cumulative survival (at 1, 2, 3, and 4 years), and major complications did not differ significantly between groups.

In another 2013 study by Ding et al, complications were retrospectively compared for 556 patients treated with MWA for 1090 tumors (491 HCC, 18 cholangiocarcinoma, 47 liver metastases) and 323 patients treated with RFA for 562 liver tumors (279 HCC, 6 cholangiocarcinoma, 38 liver metastases). (11) Rates of death (2/556 MWA, 1/323 RFA patients) as well as major and minor complications did not differ significantly between groups.

In 2013, Takami et al reported on 719 patients treated with MWA for HCC (mean tumor size, 26.9 mm) at a single institution. (12) OS rates were 97.7% at 1 year, 62.1% at 5 years, and 34.1% at 10 years. For 390 patients with 3 or fewer tumors measuring 3 cm or less, OS rates were 97.9% at 1 year, 70.0% at 5 years, and 43.0% at 10 years. When MWA results were compared with 34 patients treated at the same institution with hepatic resection, OS, DFS, and local recurrence rates did not differ significantly.

In a 2012 report on needle-track seeding, Yu et al followed 1462 patients treated with MWA for 2530 liver tumors over a 14-year period. (13) Twelve seeding nodules with a mean size of 2.3 cm (range, 1.3-3.9 cm) were found in 11 patients within 6 to 37 months (median, 10 months) after receiving MWA.

In April 2011, Simo et al retrospectively compared MWA (13 patients with 15 HCC tumors) with RFA (22 patients with 27 HCC tumors) performed by a single surgeon. (14) No significant differences were identified between treatment group characteristics, except for sex (54% vs 86% male, respectively). Average tumor size was 2.31 cm in the MWA group and 2.53 cm in the RFA group. Average tumor ablation volumes did not differ significantly for MWA (28.99 cm) and RFA (23.43 cm). In the MWA group, at a mean 7-month follow-up, DFS was 54%, with 2 patients having received liver transplants, 31% having disease progression and 15% deceased. Mean follow-up in the RFA group was 19 months. This group experienced 50% OS: 4% of patients had liver transplants, 9% had disease progression, and 36% died. Operative times were shorter in the MWA group (112 minutes vs 149 minutes)

**Case Series**

In 2011, Zhou et al. prospectively evaluated percutaneous MWA for HCC in 215 patients with tumors of 60 mm or less (median size, 29 mm) in a single center, Phase II study. (15) The authors reported technical effectiveness in all patients. OS rates at 1, 2, 3, 4, and 5 years were 94%, 82.9%, 66%, 54.1%, and 44.4%, respectively, and median survival time was 40 months (range, 4-106 months). Complications related to the procedure included 3 cases of pleural effusion and one case of bile duct injury.

In another prospective study by Zhou et al. in 2009, percutaneous MWA was performed on 124 patients with 144 HCC lesions and 28 patients with 35 lesions of hepatic metastases. (16) Included in this total of 152 patients were 59 patients with 61 lesions (mean size, 27 mm) located less than 5 mm from the gastrointestinal tract and 93 patients with 126 lesions (mean size 24 mm) located more than 5 mm from the gastrointestinal tract. For lesions less than 5 mm from the gastrointestinal tract, the temperatures of the margins were monitored closely during ablation and to prevent thermal injury, ethanol injections were placed into marginal tumor tissue in 33 lesions that were protruding or in contact with the gastrointestinal tract. No procedural complications were noted; however, tumor seeding occurred in 3 patients. Complete ablation was achieved in 47 of 53 lesions (88.7%) in the group with tumors near the gastrointestinal tract and in 116 of the other 126 lesions (92.1%), as confirmed by imaging during the follow-up period ranging from 3 to 32 months. Local tumor progression occurred in 16 tumors during 1- to 9-month follow-up. Separate treatment outcomes for hepatocellular tumors and hepatic metastasis were not provided.
In 2009, Liang et al retrospectively reviewed complications experienced with MWA for the treatment of 1928 malignant liver tumors in 1136 patients at a single institution. Each patient received an average of 1.8 treatment sessions (total treatment sessions, 3697). Thirty (2.6%) patients experienced major complications, which included 5 cases of liver abscess and empyema, 2 bile duct injuries, 2 colon perforations, 5 tumor seedings, 12 pleural effusions requiring thoracentesis, 1 hemorrhage requiring arterial embolization, and 3 skin burns requiring. Two deaths occurred within 14 days of MWA in patients with Child-Pugh class B uncompensated cirrhosis. One patient (age 78 years) had multiorgan failure and another (age 83 years) had respiratory and cardiac failure. Minor more frequent complications included fever (83.4%), pain (80.1%), asymptomatic pleural effusion (10.4%), and thickening of the gallbladder wall (2.8%), and arteriportal shunt (0.3%), small stricture of the bile duct (0.4%), and skin burn requiring no treatment (1.6). A significantly higher rate of major complications and more ablation sessions were experienced when a non-cooled-shaft antenna was used during the period of 1994 to 2005 (n=583) than with newer technology; cooled-shaft antennas were used beginning in 2005 (n=583).

Taniai et al (2006) reported on 30 patients with multiple HCC tumors who underwent reduction hepatectomy with postoperative TACE. Before surgery, patients were randomly assigned to receive no intraoperative adjuvant therapy (n=15) or intraoperative adjuvant therapy with either MWA (n=10) or RFA (n=5) of satellite lesions. No significant differences were identified between the no intraoperative adjuvant therapy and intraoperative adjuvant therapy groups, including sex, age, nodule size (maximum tumor size, 42.7±23.5 mm vs 37.8±16 mm, respectively), Child-Pugh cirrhosis class, and number of nodules. Cumulative survival rates at 3 and 5 years did not differ significantly between the no intraoperative adjuvant therapy group (35.0% and 0%, respectively) and the intraoperative adjuvant therapy group (35.7% and 7.7%, respectively). A-fetoprotein, number of tumors, maximum tumor size, and clinical stage, but not intraoperative adjuvant therapy, were identified as independent prognostic survival factors.

Lu et al (2005) reported a retrospective comparison of 102 patients with HCC treated with MWA (49 patients with 98 nodules; mean size, 2.5 cm) or RFA (53 patients with 72 nodules; mean size, 2.6 cm). Patient follow-up was about 25 months in both groups. Complete ablation did not differ significantly between groups (95% [93/98] tumors in the MWA group vs [93% [67/72] tumors in the RFA group). However, complete ablation rates improved for smaller tumors of less than 3 cm in size to 98.6% (73/74) in the MWA group and to 98% (50/51) in the RFA group. In tumors larger than 3 cm, complete ablation rates declined to 83.3% (20/24) in the MWA group and to 81% (17/21) in the RFA group. There were also no significant differences between groups in rates of local tumor recurrence (11.8% for MWA vs 20.9% for RFA), major complications (8.2% vs 5.7%, respectively), or DFS at 1, 2, and 3 years (45.9%, 26.9%, and 26.9% vs 37.2%, 20.7%, and 15.5%, respectively).

In 2002, Shibata et al reported on 72 consecutive patients with 94 small HCC nodules randomized by sealed envelope to MWA or to RFA performed by a single surgeon. No significant differences were identified between treatment group characteristics (eg, sex, age, nodule size, Child-Pugh class, number of nodules). In the RFA group, complete ablation was seen in 46 (96%) of 48 nodules (mean size, 2.3 cm; range, 1.0-3.7 cm) and 41 (89%) of 46 nodules (mean size, 2.2 cm; range, 0.9-3.4 cm) treated with MWA (p=0.26). Treatment outcomes did not differ significantly between groups in rates of untreated disease during the 6- to 27-month follow-up (8/46 nodules for MWA vs 4/48 nodules for RFA), or major complication rates (4 vs 1, respectively). Major complications included 1 case of segmental hepatic infarction in the RFA group compared to 1 case of each of the following in the MWA group: liver abscess, cholangitis with intrahepatic bile duct dilatation, subcutaneous abscess with skin burn, and subcapsular hematoma. Life-threatening complications were not reported. The number of treatment sessions required per nodule in the RFA group (1.1) was significantly lower than in the percutaneous MWA group (2.4; p<0.001). However, treatment time per session was significantly shorter with MWA (33 minutes) than with RFA (53 minutes).

**Hepatic Metastasis from Primary Cancers from Other Sites**

**Systematic Reviews**

A 2014 Health Technology Assessment and a 2013 Cochrane review reported on ablation for liver metastasis. The reviewers found insufficient evidence to determine any benefits of MWA for liver metastasis over surgical resection.

In Bertot’s 2011 systematic review (previously described), only 1 RCT was identified comparing MWA for hepatic metastases to the criterion standard of surgical resection.

In 2011, Pathak et al conducted a systematic review of ablation techniques for colorectal liver metastases, which
included 13 studies on MWA (total N=406 patients) with a minimum of 1-year follow-up. (21) Mean survival rates were 73%, 30%, and 16% and ranged from 40% to 91.4%, 0% to 57%, and 14% to 32% at the 1-, 3-, and 5-year follow-ups, respectively. Minor and major complication rates were considered acceptable, and ranged from 6.7% to 90.5% and 0% to 19%, respectively. Local recurrence rates ranged from 2% to 14%.

In the 2009 systematic review by Ong (previously described), local recurrence rates for liver metastases after MWA treatment averaged 15%, but varied between 0% and 50% in the 7 studies that addressed liver metastases. (7)

**Clinical Studies**

In 2013, Liu et al reported on liver metastases for 35 patients treated with MWA (62 tumors) and 54 patients treated with RFA (70 tumors). Ablation was complete in 89% (117/132) of tumors and did not differ significantly between tumor types: 86% (56/65) for metastatic colorectal cancer and 91% (61/67) for other metastatic disease. Tumors 3.0 cm or smaller were completely ablated significantly more often than tumors larger than 3.0 cm (94% vs 67%, p=0.001).

In 2011, Lorentzen et al retrospectively reviewed MWA in 39 patients with 125 liver metastases from the primary sites of colorectal cancer (n=31), breast cancer (n=6), carcinoid tumor (n=1), and gastrointestinal stromal tumor (n=1). (22) Complete ablation was achieved in 100% of tumors (median size, 1.5 cm) with 1 treatment session in 34 patients, in 2 sessions for 4 patients, and in 3 sessions for 1 patient. One case of liver abscess, which resolved after percutaneous drainage, was the only major complication reported. Four minor complications were reported (1 incidence of ascites, 3 complaints of puncture site pain). At median follow-up of 11 months, local tumor progression was seen in 12 (10%) of 125 tumors in 10 (26%) of the 39 patients.

In a prospective, single-institution, phase 2 study in 2010, Martin et al reported on 100 patients treated with 270 open or MWA for HCC (n=17) and liver metastases from the primary sites of colorectal (n=50), carcinoid (n=11), and other cancers (n=22, including cholangiocarcinoma, metastatic breast, renal cell carcinoma, bladder, carcinoid, melanoma, and sarcoma). (23) Median tumor size was 3.0 cm. Thirty-eight patients received MWA, 53 patients had MWA plus concomitant hepatic resection, and 9 patients had MWA concomitant with other organ resection. Only 2 patients had incomplete ablations after the procedure. No bleeding complications were experienced, but 2 cases of hepatic abscess and 2 cases of hepatic insufficiency occurred. At median follow-up of 36 months, 5 patients had incomplete ablations and 2 (2%) patients had local tumor recurrence; 37 (37%) patients developed recurrence at nonablated sites.

In 2000, Shibata et al reported on 30 patients with hepatic metastases from colorectal cancer randomly assigned without stratification to MWA after laparotomy (n=14) or to hepatectomy (n=16). (24) Of the original 40 patients, 10 patients were excluded because researchers discovered intraoperatively that they did not meet study criteria (they had extensive metastasis or ≥10 tumors). The 2 treatment groups did not significantly in age (mean age, 61 years in both groups), number of tumors (mean, 4.1 vs 3.0, respectively), or tumor size (mean, 27 mm vs 34 mm, respectively). No significant differences were observed in survival rates (27 months for MWA vs 25 months for hepatectomy) or mean DFS (11.3 months for MWA vs 13.3 months for hepatectomy). However, intraoperative blood loss was significantly lower and no blood transfusions were required in the MWA group (6 patients in the hepatectomy group required transfusions). Complications in the MWA group included 1 hepatic abscess and 1 bile duct fistula. In the hepatectomy group, complications were 1 intestinal obstruction, 1 bile duct fistula, and wound infection.

**Lung Cancer**

In 2015, Acksteiner and Steinke reported a retrospective study that evaluated the safety, effectiveness, and follow-up imaging of MWA in 10 patients (age range, ≥75 years) with early-stage non-small-cell lung cancer (NSCLC). (25) Follow-up with CT and 18-fluorodeoxyglucose-positron emission tomography (FDG-PET) extended for 30 months (median, 12 months). No periprocedural deaths or major complications were reported. Seven patients were disease-free. Three patients showed growth of the treated lesions, 1 patient died (age 90) due to unknown cause 18 months postsurgery. One patient still living presented with local progression and disseminated metastatic disease at 12 months. One patient showed increasing soft tissue mass at the ablation site 15 months posttreatment, but 3 consecutive core biopsies over 2 months failed to confirm tumor recurrence.

A 2015 observational study evaluated the clinical efficacy and utility of percutaneous microwave ablation therapy (PMAT) for lung cancer without surgical treatment. (26) Thirty-nine lesions in 29 patients with peripheral lung
cancer were treated by PMAT under local anesthesia. Treatments were completed in 29 patients. Average surgical time was 8 minutes (range, 5-12 minutes). Eight, 14, 4, and 3 patients achieved complete remission, partial remission, stable status, and progression, respectively, for an effectiveness rate of 76%. Complications included 5, 2, and 15 cases of pneumothorax, pleural effusion, and fever, respectively. No complications from needle track insertion were observed. Mean progression-free survival was 15 months. One- and 2-year OS rates were 91% and 83%, respectively.

In 2012, Lu et al retrospectively reviewed 69 patients treated with MWA for inoperable lung cancer or metastatic pulmonary metastases. (27) OS rates for patients with pulmonary metastases at 1, 2, and 3 years were 48%, 24%, and 14%, respectively. The recurrence-free survival rates for patients with NSCLC at 1, 2, and 3 years were 73%, 50%, and 27%, respectively. OS rates were 67% at 1, 45% at 2, and 25% at 3 years. Pneumothorax occurred in 25% of patients.

In 2013, Belfiore et al reported on a retrospective review of 56 patients treated with MWA for inoperable lung cancer or metastatic pulmonary metastases. (28) Disease-free survival rates were 69% at 1 year, 54% at 2 years, and 49% at 3 years. Pneumothorax was reported in 18 (32%) patients.

In 2011, Vogl et al prospectively assessed 80 patients treated with MWA for inoperable pulmonary metastases. (29) Rates were 91% at 1 year and 75% at 2 years. Pneumothorax occurred in 11 (9%) of 130 MWA sessions, and pulmonary hemorrhage occurred in 8 (6%) of 130 sessions.

**Primary Renal Tumors**

**Systematic Reviews**

In a 2014 systematic review and meta-analysis, Katsanos et al. compared thermal ablation (MWA and RFA) with surgical nephrectomy for small renal tumors (mean size 2.5 cm). (30) Included in the analysis were 1 randomized study on MWA (31) (described next) and 5 cohort studies on RFA with a total of 587 patients. In the ablation group, the complication rates and renal function decline were significantly lower than in the nephrectomy group (p=0.04 and p=0.03, respectively). The local recurrence rate was 3.6% in both groups (risk ratio=0.92, 95% CI, 0.4 to 2.14, p=0.79) and disease-free survival up to 5 years was not significantly different between groups (hazard ratio=1.04, 95% CI, 0.48 to 2.24, p=0.92).

Martin et al reported on a meta-analysis of MWA versus cryoablation for small renal tumors in 2013. (32) Included in the analysis were 7 MWA studies (n=164) and 44 cryoablation studies (n=2989). The studies were prospective or retrospective, nonrandomized, noncomparative studies. The mean follow-up duration was shorter for MWA than cryoablation (17.86 months vs 30.22 months, p=0.07). While the mean tumor size was significantly larger in the MWA studies than the cryoablation studies (2.58 cm vs 3.13 cm, respectively, p=0.04), local tumor progression (4.07% vs 2.53%, respectively; p=0.46), and progression to metastatic disease (0.8% vs 0%, respectively; p=0.12) were not significantly different.

**Clinical Studies**

In 2012, Guan et al reported on a prospective randomized study to compare the use of MWA with partial nephrectomy (the criterion standard of nephron-sparing surgical resection) for solitary renal tumors less than 4 cm. (31) Forty-eight patients received MWA and 54 had partial nephrectomy. Patients in the MWA group had significantly fewer postoperative complications than the partial nephrectomy group (6 [23.5%] vs. 18 [33.3%]; p=0.019). MWA patients also had significantly less postoperative renal function declines (p<0.009) and estimated perioperative blood loss (p<0.001) than partial nephrectomy patients. At last follow-up, estimated glomerular filtration rate declines in both groups were similar (p=1.000). Disease-specific deaths did not occur, and overall local recurrence-free survival by Kaplan-Meier estimates at 3 years were 91.3% for MWA and 96.0% for partial nephrectomy (p=0.541). Longer follow-up is needed.

Several small case studies on renal tumors have been reported. In 2012, Yu et al. reported on a retrospective review of 46 patients treated with MWA for renal cell carcinoma. (33) Complete ablation occurred in 98% of tumors (48 of 49), which had a mean tumor size of 3.0±1.5 cm. At a median follow-up of 20.1 months, all 46 patients were metastasis-free. OS rates were 100% at 1 and 2 years and 97.8% at 3 years.

In 2011, Muto et al reported on complete tumor coagulation necrosis in 10 patients treated with MWA for clear cell renal carcinoma (median tumor size, 2.75 cm). (34) Depending on tumor size, the microwave antennas were used
1 to 3 times and mean application time were 14.1 minutes. No complications were reported during or after the procedure. Bai et al (2010) reported complete laparoscopic MWA in 17 of 18 clear cell renal carcinoma tumors (mean tumor size, 2.8 cm). In this study, evidence of disease progression was not found at a median follow-up of 20 months, including a patient with incomplete ablation followed for 31 months. Complications reported were mild (18.2%), and renal function did not significantly deteriorate.

In another study, Guan et al. reported on the safety of retroperitoneoscopic MWA for renal hamartoma in 2010. In this case series report, 15 of 16 patients had complete tumor ablation. Disease recurrence was not found in all 16 patients at a median follow-up of 16 months.

Other Tumors or Conditions
No RCTs on the use of MWA for other tumors or conditions were identified. Case studies and retrospective reviews on MWA for adrenal carcinoma, metastatic bone tumors, intrahepatic primary cholangiocarcinoma, benign thyroid tumors, and other non-oncologic conditions (i.e., bleeding peptic ulcers, esophageal varices, secondary hypersplenism) were identified.

A systematic review of ablation therapies, including MWA, for locally advanced pancreatic cancer was published in 2014. The reviewers found limited available evidence on MWA for pancreatic cancer. Therefore, without randomized studies, no conclusions could be drawn on thermal ablation methods for pancreatic cancer.

Ongoing and Unpublished Clinical Trials
Some currently unpublished trials that might influence this review are listed in Table 1

<table>
<thead>
<tr>
<th>NCT No.</th>
<th>Trial Name</th>
<th>Planned Enrollment</th>
<th>Completion Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ongoing</td>
<td>NCT01340105 Microwave Versus Radiofrequency Ablation for Hepatocellular Carcinoma: a Prospective Randomized Control Trial</td>
<td>92</td>
<td>Apr 2016</td>
</tr>
</tbody>
</table>

NCT: national clinical trial.

Summary of Evidence
The evidence for microwave ablation (MWA) in individuals who have an unresectable primary or metastatic tumor (eg, hepatic [primary or metastatic], pulmonary, renal) includes case series, observational studies, cohort studies, randomized controlled trials (RCTs), and systematic reviews. Relevant outcomes are overall survival, disease-specific survival, symptoms, quality of life, and treatment-related mortality and morbidity. Available studies have shown that MWA results in a wide range of complete tissue ablation (50%-100%) depending on tumor size, with complete ablation common and nearing 100% with smaller tumors (eg, less than or equal to 3 cm). Tumor recurrence rates at ablated sites are very low. However, tumor recurrence at nonablated sites is common and may correlate with disease state (eg, in hepatocellular carcinoma). Intraoperative and postoperative minor and major complications are low, especially when tumors are smaller and accessible. Patient selection criteria and rationale for using MWA over other established techniques (eg, surgical resection, radiofrequency ablation) are needed. The evidence is insufficient to determine the effects of the technology on health outcomes.

Clinical Input Received from Physician Specialty Societies and Academic Medical Centers
While the various physician specialty societies and academic medical centers may collaborate with and make recommendations during this process through the provision of appropriate reviewers, input received does not represent an endorsement or position statement by the physician specialty societies or academic medical centers, unless otherwise noted.

2015
In response to requests, input was received from 2 physician specialty societies and 1 academic medical center while this policy was under in 2015. This number of responses was less than optimal. Input overall was mixed. There was some support for the medical necessity of MWA in each category, with some reviewers indicating that it was standard of care for certain tumors. However, there were no indications for which all 3 reviewers agreed that MWA should be medically necessary.
In response to requests, input was received from 2 physician specialty societies (3 reviews) and 4 academic medical centers (6 reviews) while this policy was in development. Eight reviewers considered MWA investigational to treat primary tumors such as hepatocellular carcinoma (HCC), benign and malignant renal tumors, lung tumors, adrenal tumors, or cholangiocarcinoma. The reviewers noted insufficient evidence and a need for further studies on MWA. However, 1 reviewer indicated microwave ablation for primary tumors, including, but not limited to HCC, benign and malignant renal tumors, lung tumors, adrenal tumors and cholangiocarcinoma, may be considered a treatment option, and another reviewer indicated that MWA for renal tumors may be considered a treatment option.

Four reviewers considered MWA investigational to treat liver metastases. However, 2 reviewers indicated MWA for liver metastases may be considered a treatment option. One reviewer noted MWA may be appropriate for tumors not amenable to radiofrequency ablation (RFA) or other local treatments. This reviewer also indicated microwave ablation may be more appropriate for tumors located near large blood vessels.

Practice Guidelines and Position Statements

National Comprehensive Cancer Network (NCCN)
The NCCN guidelines on hepatobiliary cancers lists MWA (along with RFA, cryoablation, and percutaneous alcohol injection) as a treatment option for HCC tumors in patients who are not candidates for potential curative treatments (e.g., resection and transplantation) and do not have large-volume extrahepatic disease. (43) Ablation should only be considered when tumors are accessible by percutaneous, laparoscopic, or open approaches The guidelines indicate HCC tumors of 3 centimeters or less may be curatively treated with ablation alone. HCC tumors between 3 and 5 centimeters may also be treated with ablation to prolong survival when used in combination with arterial embolization. Additionally, the tumor location must be accessible to permit ablation of the tumor and tumor margins without ablating major vessels, bile ducts, the diaphragm or other abdominal organs. However, there are only 2 reviews cited in the guideline on ablative techniques to support these recommendations, but these reviews are not specific to MWA [category 2A].

NCCN guidelines for neuroendocrine tumors, (v.1.2015) do not mention MWA. (44) Guidelines state that: “Cytoreductive surgery or ablative therapies such as radiofrequency ablation (RFA) or cryoablation may be considered if near-complete treatment of tumor burden can be achieved (category 2B). For unresectable liver metastases, hepatic regional therapy (arterial embolization, chemoembolization, or radioembolization [category 2B]) is recommended.”

National Institute for Health and Clinical Excellence (NICE)
The NICE published updated guidance on MWA for the Treatment of Metastases in the Liver in August 2011. (45) This guidance indicates “Current evidence on microwave ablation for the treatment of liver metastases raises no major safety concerns. The evidence on efficacy is inadequate in quantity and quality. Therefore this procedure should only be used with special arrangements for clinical governance, consent and audit or research.” NICE also published guidance on MWA for HCC in 2007. (46) This guidance indicated “Current evidence on the safety and efficacy of microwave ablation of hepatocellular carcinoma appears adequate to support the use of this procedure provided that the normal arrangements are in place for consent, audit and clinical governance.” The guidance also stated there are no major concerns about the efficacy of MWA but noted there is limited long-term survival data available.

American College of Chest Physicians (CHEST)
The 2013 evidence-based guidelines from CHEST on the treatment of non-small cell lung cancer (NSCLC) note that the role of ablative therapies in the treatment of high-risk patients with stage I NSCLC is evolving. (46) RFA, the most studied of the ablative modalities, has been used effectively in medically inoperable patients with small (less than 3 cm) peripheral NSCLC that are clinical stage I.

U.S. Preventive Services Task Force Recommendations
N/A
Medicare National Coverage
There is no national coverage determination (NCD). In the absence of an NCD, coverage decisions are left to the discretion of local Medicare carriers.

References


<table>
<thead>
<tr>
<th>Date</th>
<th>Reason</th>
</tr>
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<tbody>
<tr>
<td>02/27/12</td>
<td>New Policy – Add to Surgery section. Policy created with literature review through October 2011; investigational for all tumors.</td>
</tr>
<tr>
<td>07/31/12</td>
<td>Code 47379 added to the policy as this procedure can be performed laparoscopically</td>
</tr>
<tr>
<td>09/07/12</td>
<td>Update coding section – ICD-10 codes are now effective 10/01/14.</td>
</tr>
<tr>
<td>12/20/12</td>
<td>Update Related Policies; policy number 7.01.540 was replaced with 7.01.95.</td>
</tr>
<tr>
<td>04/16/13</td>
<td>Replace policy. Policy updated with literature review; reference numbers 2, 12-13, 21-25, 32 and 36 added. Policy statement unchanged.</td>
</tr>
<tr>
<td>03/11/14</td>
<td>Coding Update. Code 55.33 was removed per ICD-10 mapping project; this code is not utilized for adjudication of policy.</td>
</tr>
<tr>
<td>12/08/15</td>
<td>Annual Review. Policy updated with literature search; no change to the policy statement.</td>
</tr>
<tr>
<td>05/10/16</td>
<td>Annual Review. Policy updated with literature review through February 15, 2016; references added. Clinical input added. Policy statement unchanged. CPT code 0301T added to this policy.</td>
</tr>
</tbody>
</table>

Disclaimer: This medical policy is a guide in evaluating the medical necessity of a particular service or treatment. The Company adopts policies after careful review of published peer-reviewed scientific literature, national guidelines and local standards of practice. Since medical technology is constantly changing, the Company reserves the right to review and update policies as appropriate. Member contracts differ in their benefits. Always consult the member benefit booklet or contact a member service representative to determine coverage for a specific medical service or supply. CPT codes, descriptions and materials are copyrighted by the American Medical Association (AMA). ©2016 Premera All Rights Reserved.
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U.S. Department of Health and Human Services
200 Independence Avenue SW, Room 509F, HHH Building
Washington, D.C. 20201, 1-800-368-1019, 800-537-7697 (TDD)
Complaint forms are available at https://ocrportal.hhs.gov/ocr/portal/lobby.jsf or by mail at:

Please call 800-368-1019, 1-800-537-7697 (TDD), or TTY 800-842-5357.

Getting Help in Other Languages

This Notice has Important Information. This notice may have important information about your application or coverage through Premera Blue Cross. There may be key dates in this notice. You may need to take action by certain deadlines to keep your health coverage or help with costs. You have the right to get this information and help in your language at no cost. Call 800-722-1471 (TTY: 800-842-5357).

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本通知有重要的訊息。本通知可能有關於您透過 Premera Blue Cross 提交的申請或保險的重要訊息。本通知內可能有重要日期，您可能需要在截止日期之前採取行動，以保留您的健康保險或費用補貼。您有權利免費以您的母語得到本訊息和幫助。請接電話 800-722-1471 (TTY: 800-842-5357)。

Oromo (Cushite):


Français (French):


Kreyòl ayisyen (Creole):


Deutsche (German):


Hmoob (Hmong):


Ilokano (Ilocano):

Daytoy a Pakdaar ket naglaon iti Napateg nga Impormasion. Daytoy a pakdaar mabaliti nga adda ket naglaon iti napateg nga impormasion maipanggpe iti aplikasyonu woyen coverage babaen iti Premera Blue Cross. Daytoy ket mabaliti dagiti importante a pelta iti daytoy a pakdaar. Mabaliti nga adda rumbeng nga aramidengu nga adda sangbay dagitii partikular a naituling nga adda aldaw tapo napagtagalnediytoy nga coverage ti salun-atyo woyen tulong kadaagi gastos. Adda karbenganyo a mangala iti daytoy nga impormasion ken tulong ti bukodyo a pagasasan nga awan ti bayadanyo. Tumawag ti numero nga 800-722-1471 (TTY: 800-842-5357).

Italiano (Italian):

Premera Blue Cross

Determine when to seek help for medical care if you have
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