Repetitive Transcranial Magnetic Stimulation (rTMS)

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**Coverage:**

**Initial rTMS Treatment**

Repetitive transcranial magnetic stimulation (rTMS) **may be considered medically necessary** in the acute phase treatment* of major depression, single episode or recurrent, ONLY if ALL of the following conditions are met:

1. Diagnosis of major depression, either single episode or recurrent (non-psychotic); AND

2. Patient has had 4 failed trials of antidepressant medications from at least 2 different classes of antidepressants in the current episode; AND

3. Patient is currently, or has been, in formal cognitive behavioral therapy; AND

4. National standardized rating scales such as PHQ-9 are administered weekly during treatment;

5. None of the following conditions or contraindications to rTMS are present:
   a. Seizure disorder or any history or seizure disorder (except those induced by ECT or isolated febrile seizures in infancy without subsequent treatment or recurrence); OR
   b. Presence of acute or chronic psychotic symptoms or disorders (e.g., schizophrenia, schizophrainiform or schizoaffective disorder) in the current depressive episode; OR
c. Neurological conditions that include epilepsy, cerebrovascular disease, dementia, increased intracranial pressure, history of repetitive or severe head trauma, or primary or secondary tumors in the central nervous system; OR
d. Excessive use of alcohol or illicit substances within the last 30 days;
e. The patient did not respond to a prior course of rTMS treatments (as defined by not achieving at least a 50% reduction in severity of scores for depression in a standardized rating scale such as the PHQ-9 by the end of acute phase treatment, i.e., 36 sessions).

*NOTE: If the above conditions are met, 36 acute phase sessions (30 sessions intended to be given over 6 weeks, plus 6 taper-down sessions) of rTMS can be authorized.

Subsequent rTMS Treatments

Subsequent acute phase treatments for rTMS may be considered medically necessary if the patient has met all of the following conditions:

1. Patient has documented positive response to prior rTMS treatment, as defined by at least a 50% reduction in severity of scores for depression on a standardized rating scale such as the PHQ-9 by the end of acute phase treatment; AND

2. Patient has not received a separate acute phase rTMS treatment within the last 6 months; AND

3. Patient has none of the following conditions or contraindications to rTMS:

   a. Seizure disorder or any history or seizure disorder (except those induced by ECT or isolated febrile seizures in infancy without subsequent treatment or recurrence); OR

   b. Presence of acute or chronic psychotic symptoms or disorders (e.g., schizophrenia, schizophreniform or schizoaffective disorder) in the current depressive episode; OR

   c. Neurological conditions that include epilepsy, cerebrovascular disease, dementia, increased intracranial pressure, history of repetitive or severe head trauma, or primary or secondary tumors in the central nervous system; OR
d. Excessive use of alcohol or illicit substances within the last 30 days.

Subsequent repetitive transcranial magnetic stimulation (rTMS) is considered experimental, investigational, and/or unproven for:

1. A patient who did not respond to a prior episode of rTMS treatments (as defined by not achieving at least a 50% reduction in severity of scores for depression on a standardized rating scale such as the PHQ-9 by the end of acute phase treatment, i.e., 36 sessions);

2. Maintenance treatment of major depression (i.e., “booster treatments”).

Other Circumstances for rTMS Treatment

Repetitive transcranial magnetic stimulation (rTMS) is considered experimental, investigational, and/or unproven in all other circumstances, including but not limited to:

1. The patient is actively psychotic;

2. The patient has dementia or a cognitive disorder;

3. The patient has excessive use of alcohol or illicit substances within the last 30 days;

4. Any other psychiatric or neurologic disorder including, but not limited to:
   a. Schizophrenia;
   b. Migraine headaches;
   c. Epilepsy or other seizure disorder, or any history or seizure disorder (except those induced by ECT or isolated febrile seizures in infancy without subsequent treatment or recurrence);
   d. Cardiovascular disease/stroke;
   e. Dementia;
   f. Alzheimer’s disease;
   g. Attention deficit disorder/hyperactivity disorder;
   h. Bulemia nervosa;
   i. Dysphagia;
   j. Fibromyalgia;
Transcranial magnetic stimulation (TMS) is a noninvasive method of delivering electrical stimulation to the brain. A magnetic field is delivered through the skull where it induces electric currents that affect neuronal function. Repetitive TMS (rTMS) is being evaluated as a treatment of depression and other psychiatric/neurologic brain disorders.

**Background**

Transcranial magnetic stimulation (TMS) was first introduced in 1985 as a new method of noninvasive stimulation of the brain. The technique involves placement of a small coil over the scalp; passing a rapidly alternating current through the coil wire, which produces a magnetic field that passes unimpeded through the scalp and bone, resulting in electrical stimulation of the cortex. TMS was initially used to investigate nerve conduction; for example, TMS over the motor cortex will produce a contralateral muscular-evoked potential. The motor threshold, which is the minimum intensity of stimulation required to induce a motor response, is empirically determined for each individual by localizing the site on the scalp for optimal stimulation of a hand muscle, then gradually increasing the intensity of stimulation. The stimulation site for treatment of depression is usually 5 cm anterior to the motor stimulation site.

Interest in the use of TMS as a treatment for depression was augmented by the development of a device that could deliver rapid, repetitive stimulation. Imaging studies had shown a decrease in activity of the left dorsolateral prefrontal cortex (DLPFC) in depressed patients, and early studies suggested that high frequency (e.g., 5–10 Hz) TMS of the left DLPFC had antidepressant effects. Low frequency (1–2 Hz) stimulation of the right DLPFC has also been investigated. The rationale for low frequency TMS is inhibition of right frontal cortical activity to correct the interhemispheric imbalance. A combination approach (bilateral stimulation), or deep stimulation with an H1 coil, are also being explored. In contrast to electroconvulsive therapy, TMS does not require anesthesia and does not induce a convulsion.
rTMS is also being tested as a treatment for a variety of other disorders including alcohol dependence, Alzheimer’s disease, neuropathic pain, obsessive-compulsive disorder (OCD), post-partum depression, Parkinson disease, stroke, posttraumatic stress disorder, panic disorder, epilepsy, dysphagia, Tourette’s syndrome, schizophrenia, migraine, spinal cord injury, fibromyalgia, and tinnitus. In addition to the potential for altering interhemispheric imbalance, it has been proposed that high frequency rTMS may facilitate neuroplasticity.

Regulatory Status

Devices for transcranial stimulation have received clearance by the U.S. Food and Drug Administration (FDA) for diagnostic uses. One device, NeoPulse (Neuronetics, Atlanta, GA), received approval as a therapy for depression. Initially examined by the FDA under a traditional 510(k) application, the NeoPulse, now known as NeuroStar® TMS, received clearance for marketing as a “De Novo” device in 2008. In March 2014, NeuroStar® TMS received FDA 510(k) approval for use in adult patients who have failed to receive satisfactory improvement from prior antidepressant medication in the current episode. The Brainsway H-Coil Deep TMS device (Brainsway Ltd.) received FDA clearance in 2013. This device is indicated for the treatment of depression in patients who have failed to respond to antidepressant medications in their current episode of depression.

Note: An FDA advisory panel met in January 2007 to determine if the risk-to-benefit profile for the NeoPulse was comparable to the risk-to-benefit profile of predicate electroconvulsive therapy (ECT) devices. The panel was not asked for a recommendation regarding the regulatory determination of substantial equivalence for this 510(k) submission. Materials presented at the Neurological Devices Panel meeting, as well as a summary, are posted online at: <www.fda.gov>.

Rationale:

The Blue Cross and Blue Shield Association (BCBSA) Technology Evaluation Center (TEC) published an assessment of repetitive TMS (rTMS) for depression in 2009. The TEC Assessment concluded that the available evidence did not permit conclusions regarding the effect of TMS on health outcomes. Limitations of the evidence included:

- Equivocal efficacy in the largest sham-controlled trial of TMS,
- Uncertain clinical significance of the short-term anti-depressant effects found in meta-analyses,
• A lack of information beyond the acute period of treatment, and
• Lack of comparison with standard therapy (a second course of antidepressant therapy) in the population for whom TMS is indicated (patients who have failed one 6-week course of antidepressant medication).

BCBSA published an updated TEC Assessment of TMS for depression in 2011. Included were six recent meta-analyses, the largest of which evaluated 30 double-blind sham-controlled trials with a total of 1,383 patients. Recent clinical trials were also reviewed. The 2011 TEC Assessment reached the following conclusions:

• The meta-analyses and recent clinical trials of TMS generally show statistically significant effects on depression outcomes at the end of the TMS treatment period. However, the clinical significance and durability of the effect are not well-characterized.
• The largest randomized clinical trial showed a greater effect in patients with only one prior treatment failure, with possibly minimal or no effect in patients with greater than one prior treatment failure. Based on current evidence, it cannot be determined whether TMS after one treatment failure would be as effective as the current standard of a second course of antidepressant therapy.
• Also identified as gaps in current knowledge are whether TMS is effective as an adjunctive treatment and whether retreatment is effective.

Following is a summary of the key literature to date, focusing on randomized controlled trials (RCTs). The evidence review is divided by indication and by key differences in treatment protocols, specifically high-frequency left dorsolateral prefrontal cortex stimulation (DLPFC), low-frequency (1–2 Hz) stimulation of the right dorsolateral prefrontal cortex, or combined high-frequency and low-frequency stimulation.

**Depression**

Studies published prior to 2008 are included if the study design was a randomized sham-controlled double-blind trial that enrolled at least 40 subjects; refer to the 2008 meta-analysis by Schutter for a summary of study characteristics and stimulation parameters used in these trials. (3) Note that over the last decade, there has been a trend to increase the intensity, trains of pulses, total pulses per session, and number of sessions. (4) Unless otherwise indicated in the trials described below, stimulation was set at
100% to 120% of motor threshold, clinical response was defined as an improvement of 50% or more on the Hamilton Depression Rating Scale (HAM-D), and remission was considered to be a score of 7 or less on the HAM-D.

High Frequency rTMS of the Left Dorsolateral Prefrontal Cortex for Treatment-Resistant Depression

Lam and colleagues conducted a meta-analysis of 24 randomized controlled trials (RCTs) comparing active versus sham repetitive TMS (rTMS) in patients with TRD, although there were varying definitions of TRD. (5) This analysis calculated a number needed to treat of 6, with a clinical response in 25% of active rTMS and 9% of sham rTMS patients. Remission was reported for 17% of active rTMS and 6% of sham rTMS patients.

The largest study (23 study sites) included in the meta-analysis was a double-blind multicenter trial with 325 TRD patients randomly assigned to daily sessions of high-frequency active or sham rTMS (Monday to Friday for 6 weeks) of the left dorsolateral prefrontal cortex (DLPFC). (6) Treatment-resistant depression was defined as failure of at least 1 adequate course of antidepressant treatment. Patients had failed an average of 1.6 treatments in the current episode, with approximately half of the study population failing to benefit from at least 2 treatments. Loss to follow-up was similar in the 2 groups, with 301 (92.6%) patients completing at least 1 post-baseline assessment and an additional 8% of patients from both groups dropping out before the 4-week assessment. Intent-to-treat (ITT) analysis showed a trend favoring the active rTMS group in the primary outcome measure (2 points on the Montgomery-Asberg Depression Rating Scale (MADRS); p=0.057) and a modest (2-point) but significant improvement over sham treatment on the HAM-D. The authors reported that after 6 weeks of treatment, subjects in the active rTMS group were more likely to have achieved remission than the sham controls (14% vs. 5%, respectively), although this finding is limited by loss to follow-up.

In 2010, George et al. reported a randomized sham-controlled trial that involved 199 patients treated with left-prefrontal rTMS. (7) This was a multi-centered study involving patients with a moderate level of treatment resistance. The response rate using an ITT analysis was 14% for rTMS and 5% for sham (p=0.02). In this study, the site for stimulation was determined through pre-treatment magnetic resonance imaging (MRI). Results from Phase 2 (open treatment of non-responders) and Phase 3 (maintenance and follow-up) will be reported in the future.
Another randomized sham-controlled double-blind trial was conducted in 68 patients who had failed at least 2 courses of antidepressants. (8) Three patients in each group did not complete the 15 treatment sessions or were excluded due to a change in medication during treatment, resulting in 91% follow-up. Independent raters found a clinical response in 31% (11 of 35) of the active rTMS patients and 6% (2 of 33) of the sham group. The average change in HAM-D was 7.8 for the active group and 3.7 for the control group. The Beck Depression Inventory (BDI) decreased by 11.3 points in the active rTMS group and 4.8 points in controls. Remission was observed in 7 patients (20%) in the active rTMS group and 1 patient (3%) in the control group.

Regarding effectiveness of blinding; 15% of subjects in each group guessed that they were receiving active TMS after the first session. After the 15th session, 58% of the rTMS group and 43% of the sham group guessed that they had received active TMS; responders were more likely than non-responders (85% vs. 42%, respectively) to think that they had received the active treatment. The 11 responders were treated with antidepressant medication and followed up for 6 months. Of these, 1 was lost to follow-up, 5 (45%) relapsed, and 5 (45%) did not relapse.

Rossini and colleagues randomly assigned 54 patients who had failed at least 2 adequate courses of antidepressants to sham control or active rTMS at 80% or 100% of motor threshold (MT) for 10 sessions over a 2-week period. (9) Double-blind evaluation found an intensity-dependent response with 6% (1 of 16) of the sham, 28% (5 of 18) of the 80% MT, and 61% (11 of 18) of the 100% MT groups showing improvement of 50% or more over a 5-week evaluation. All of the patients reported that they were unaware of the differences between sham and active stimulation.

In a 2008 report, Mogg et al. randomly assigned 59 patients with major depression who had failed at least 1 course of pharmacotherapy for the index depressive episode. (10) In this study population, 78% of the patients had failed 2 treatment courses and 53% had failed 3. The sham coil, which was provided by Magstim, was visually identical to the real coil and made the same clicking sound but did not deliver a magnetic field to scalp or cortex. Blinded assessments were performed 2 days after the fifth and final (tenth) sessions (97% follow-up), with additional assessments at 6 weeks (90% follow-up) and 4 months (83% follow-up). The mean group difference was estimated to be 0.3 points in HAM-D scores for the overall analysis. Interpretation of this finding is limited, since 7 sham patients (23%) were given a course of real rTMS after the 6-week assessment and analyzed
as part of the sham group in the ITT analysis. The study was powered to detect a difference of 3.5 points in the HAM-D between the active and sham groups, and the 2.9-point group difference observed at the end of treatment was not significant. A higher percentage of patients in the active rTMS group achieved remission criteria of 8 points or less on the HAM-D (25% vs. 10% control, respectively), and there was a trend for more patients to achieve clinical response in the active rTMS group (32% vs. 10%, respectively, p=0.06). All of the 12 patients who met the criterion for clinical response (9 active and 3 sham) thought that they had received real rTMS, with more patients in the active group (70%) than the sham group (38%) guessing that they had received the real treatment. Interpretation of this finding is also limited, since the reason the subjects guessed that they had active treatment was not reported, and the subjects were not asked to guess before they began to show a clinical response.

A small double-blind randomized trial from 2009 suggests that specific targeting of Brodmann areas 9 and 46 may enhance the anti-depressant response compared with the standard targeting procedure, i.e., measuring 5 cm anterior from the motor cortex. (11) Fifty-one patients who had failed at least two 6-week courses of antidepressant medication (average 5.7 failed courses) were randomly assigned to a standard localization procedure or to structural magnetic resonance imaging (MRI)-aided localization for 3 weeks (with 1-week extension if >25% reduction on the MADRS). Six patients in the targeted group and 10 in the standard group withdrew due to lack of response. A single patient in the targeted group and 5 in the standard group withdrew for other reasons, resulting in 17 patients in the targeted group and 12 in the standard group continuing for the full 4 weeks of treatment. To adjust for the imbalance in discontinuation rates, a mixed model statistical analysis was used. There was a significant difference between the groups in the overall mixed model analysis, and planned comparisons showed significant improvement in MADRS scores for the targeted group at 4 weeks. Response criteria were met by 42% of the targeted group and 18% of the standard group. Remission criteria were met by 30% of the targeted group and 11% of the standard group. Although encouraging, additional trials with a larger number of subjects are needed to evaluate this procedure.

Several studies have compared the outcomes of rTMS with those from electroconvulsive therapy. In one study, 40 patients with nonpsychotic major depression were treated over the course of 1 month (20 total sessions) and evaluated with the HAM-D, in which
a response was defined as a 50% decrease with a final score of less than or equal to 10. (12) There was no difference in response rate between the 2 groups; 12 of 20 responded in the electroconvulsive therapy group compared to 11 of 20 in the magnetic stimulation group. A United Kingdom National Institute for Health Research health technology assessment compared efficacy and cost effectiveness of rTMS and electroconvulsive therapy. (13) Forty-six patients who had been referred for electroconvulsive therapy were randomly assigned to either electroconvulsive therapy (average of 6.3 sessions) or a 15-day course (5 treatments per week) of rTMS of the left DLPFC. Electroconvulsive therapy resulted in a 14-point improvement in the HAM-D and a 59% remission rate. Repetitive TMS was less effective than electroconvulsive therapy (5-point improvement in HAM-D and a 17% remission rate). Another study reported no significant difference between electroconvulsive therapy and rTMS in 42 patients with TRD; however, response rates for both groups were low. (14) The number of remissions (score of 7 or less on the HAM-D) totaled 3 (20%) for electroconvulsive therapy and 2 (10%) for rTMS.

Low Frequency rTMS of the Right Dorsolateral Prefrontal Cortex or Bilateral Stimulation for Treatment-Resistant Depression

Fitzgerald et al. randomly assigned 60 patients who had failed a minimum of at least two 6-week courses of antidepressant medications into 1 of 3 groups; high frequency left rTMS, low frequency right rTMS, or sham stimulation over 10 sessions. (15) All patients who entered the study completed the double-blind randomized phase, which showed no difference between the 2 active treatments (left: 13.5% reduction; right: 15% reduction) and greater improvements in the MADRS scores compared to the sham group (0.76% reduction). Only 1 patient achieved 50% improvement during the initial 2 weeks. Then, only the subjects who showed at least 20% improvement at the end of the 10 sessions (15 active and 2 sham) continued treatment. Patients who did not respond by at least 20% were switched to a different active treatment. From week 2 to week 4, there was greater improvement in the low frequency right rTMS group compared with the high frequency left rTMS group (39% vs. 14% improvement in MADRS, respectively). Seven patients (18% of 40) showed a clinical response of greater than 50% by the end of the 4 weeks.

In a subsequent study, Fitzgerald and colleagues randomly assigned 50 patients with TRD to sequential bilateral active or sham rTMS. (16) After 2 weeks of treatment, 3 subjects had
dropped out of the sham treatment group, and there was a slight but non-significant improvement favoring the active group for the MADRS (26.2 vs. 30.9, respectively) and the BDI (18.3 vs. 21.6, respectively). At this time point, 60% of subjects receiving active rTMS and 50% of subjects receiving sham treatment guessed that they were in the active group. The clinical response was reported by subjects as the major reason for their guess, with 11 of 13 responders (9 active and 2 sham) guessing that they were in the active group. As in the earlier study, only the subjects who showed at least 20% improvement at the end of each week continued treatment. Treatment on week 3 was continued for 15 subjects in the active group and 7 subjects in the sham group. By week 6, 11 subjects in the active rTMS remained in the study, with no control subjects remaining. Final ratings for the 11 subjects who continued to respond through week 6 were 8.9 on the MADRS and 9.2 on the BDI.

Another multicenter double-blind trial randomly assigned 130 patients with TRD to 5 sessions per week of either 1- or 2-Hz rTMS over the right dorsolateral prefrontal cortex. (17) Sixty-eight patients (52%) completed 4 weeks of treatment; there was an approximate 30% improvement in depression scales, with no differences between the 1- or 2-Hz groups. Due to the potential for placebo effects for this type of intervention, the absence of a sham control group limits interpretation.

A small randomized, sham-controlled trial was published in 2010 that involved either right or left rTMS in 48 patients with TRD. (18) Overall reductions in the HAM-D-24 from baseline to 3 months were not significantly different between rTMS and sham treatment groups. In this small study, right cranial stimulation was significantly more effective than left cranial stimulation (sham or rTMS).

rTMS as an Adjunctive Treatment for Moderate to Severe Depression

Schutter conducted a meta-analysis of 30 double-blind randomized sham-controlled trials (1,164 patients) of high-frequency rTMS over the left dorsolateral prefrontal cortex in patients with major depression. (3) The pooled weighted mean effect size for treatment was calculated with Hedges g, a standardized mean difference that adjusts for sampling variance, to be 0.39 (95% confidence interval [CI]: 0.25–0.54), which is considered moderate. For 27% of the population, rTMS was used as a primary/adjunctive treatment; 3 trials were included that used rTMS as a primary/adjunctive treatment for depression and
enrolled more than 40 subjects. (19-21) Repetitive TMS has also been examined in patients with clinical evidence of cerebrovascular disease and late-life depression. (22)

A 2012 study examined the efficacy of ultra-high-frequency (30 Hz) rTMS over the left prefrontal cortex in moderate to severely depressed patients who were taking medication. (23) Sham treatment consisted of low frequency stimulation to the left prefrontal cortex. No benefit of rTMS for depressive symptoms was found when lithium was added as a covariate. Ultra-high-frequency rTMS was found to improve performance on the trail-making test, which covaried with improvement of psychomotor retardation.

Additional research on whether adjunctive rTMS can improve response to pharmacologic treatment as a first-line therapy is needed.

In 2012, Carpenter et al. (58) published a multisite observational study of acute treatment outcomes of use of rTMS in real-world clinical practice settings. Forty-two US-based clinical TMS practice sites treated 307 outpatients with Major Depressive Disorder (MDD), and persistent symptoms despite antidepressant pharmacotherapy. Treatment was based on the labeled procedures of the approved TMS device. Assessments were performed at baseline, week 2, at the point of maximal acute benefit, and at week 6 when the acute course extended beyond 6 weeks. The primary outcome was change in the Clinician Global Impressions-Severity of Illness from baseline to end of acute phase. Secondary outcomes were change in continuous and categorical outcomes on self-report depression scales (9-Item Patient Health Questionnaire [PHQ-9], and Inventory of Depressive Symptoms-Self Report [IDS-SR]). Patients had a mean ± SD age of 48.6 ± 14.2 years and 66.8% were female. Patients received an average of 2.5 (± 2.4) antidepressant treatments of adequate dose and duration without satisfactory improvement in this episode. There was a significant change in CGI-S from baseline to end of treatment (-1.9 ± 1.4, P < .0001). Clinician-assessed response rate (CGI-S) was 58.0% and remission rate was 37.1%. Patient-reported response rate ranged from 56.4 to 41.5% and remission rate ranged from 28.7 to 26.5%, (PHQ-9 and IDS-SR, respectively). The authors concluded that outcomes demonstrated response and adherence rates similar to research populations, and felt the data indicate that TMS is an effective treatment for those unable to benefit from initial antidepressant medication.
Maintenance Therapy

Fitzgerald et al. reported a prospective open-label trial of clustered maintenance rTMS for patients with refractory depression. (24) All patients had received a second successful course of rTMS following relapse and were then treated with monthly maintenance therapy consisting of 5 rTMS treatments over a 2.5-day period (Friday evening, Saturday and Sunday). Patients were treated with maintenance therapy of the same type that they had initially received (14 high frequency to the left dorsolateral prefrontal cortex, 12 low frequency to the right dorsolateral prefrontal cortex, and 9 bilateral). The primary outcome was the mean duration until clinical relapse, addition or change of antidepressant medication, or withdrawal from maintenance treatment to pursue other treatment options. Of 35 patients, 25 (71%) relapsed at a mean of 10.2 months (range, 2 to 48 months), which was substantially longer than the interval (<3 months) for relapse from the initial treatment.

Janicak and colleagues reported on assessment of relapse during a multisite, open-label study. (25) In this study, patients who met criteria for partial response during either a sham–controlled or open-label phase of a prior study were tapered from rTMS and simultaneously started on maintenance antidepressant monotherapy. They were then followed for 24 weeks. Ten of 99 patients relapsed. Thirty-eight patients had symptom worsening, and 32 of these (84%) had symptomatic benefit with adjunctive rTMS.

A retrospective study that included maintenance rTMS was reported by Connolly et al. in 2012. (26) Out of the first 100 cases treated at their institution, 42 received maintenance rTMS. Most of the patients had failed more than 1 adequate antidepressant trial and were treated with high-frequency rTMS over the dorsolateral prefrontal cortex. Low-frequency rTMS to the right dorsolateral prefrontal cortex was given in patients with a family or personal history of seizures and in some patients who were also receiving high-frequency rTMS. The response rate was 50.6% of the first 100 cases and the remission rate was 24.7%. Maintenance treatment (42 patients) was tapered gradually from 2 sessions per week for the first 3 weeks to monthly. At 6 months after the initial rTMS treatment, 26 of the 42 patients (62%) maintained their response.

To date, the evidence for maintenance phase rTMS treatment for patients with MDD is limited by studies with small sample size and lack of true control groups. Furthermore, no specific guidelines for
delivering maintenance treatment currently exist, as the delivery paradigms for maintenance rTMS treatment varies across studies; additional data are needed related to durability of effect and to maintenance phases.

**Alzheimer’s Disease**

Ahmed et al. randomized 45 patients with probable Alzheimer’s disease to 5 sessions of bi-lateral high-frequency rTMS, bi-lateral low-frequency rTMS, or sham TMS over the dorsolateral prefrontal cortex. (27) Thirty-two patients had mild to moderate dementia and 13 had severe dementia. There were no significant differences between groups at baseline. Measures of cortical excitability immediately after the last treatment session showed that treatment with high-frequency rTMS reduced the duration of transcallosal inhibition. At 3 months after treatment, the high-frequency rTMS group improved significantly more than the other 2 groups in standard rating scales, and subgroup analysis showed that this was due primarily to improvements in patients with mild/moderate dementia. Patients in the subgroup of mild to moderate dementia who were treated with high-frequency rTMS improved from 18.4 to 22.6 on the Mini Mental State Examination (MMSE), from 20.1 to 24.7 on the Instrumental Daily Living Activity (IADL) scale and from 5.9 to 2.6 on the Geriatric Depression Scale (GDS).

Rabey et al. reported an industry-sponsored randomized double-blind trial of rTMS with cognitive training (NeuroAD system) in 15 patients with probable mild to moderate Alzheimer’s disease. (28) Patients received 5 sessions per week for 6 weeks over 6 different brain areas, followed by biweekly sessions for 3 months. Specific cognitive tasks were designed for the 6 targeted brain regions. These included syntax and grammar for Broca’s area, comprehension and categorization for Wernicke’s area, action naming, object naming and spatial memory tasks for the right and left dorsolateral prefrontal cortex, and spatial attention tasks for the right and left somatosensory association cortex. After 6 weeks of treatment, there was an improvement in the average Alzheimer Disease Assessment Scale, cognitive subsection (ADAS-cog) score of 3.76 points in the rTMS group compared to 0.47 in the placebo group. After 4.5 months of treatment, the ADAS-cog score in the rTMS group had improved by 3.52 points compared to a worsening of 0.38 in the placebo group. The Clinical Global Impression of Change improved significantly by an average of 3.57 after 6 weeks and 3.67 after 4.5 months compared to 4.25 and 4.29, respectively, in the placebo group.
Attention-Deficit/Hyperactivity Disorder

In 2012, Weaver et al. reported a randomized sham-controlled crossover study of rTMS in 9 adolescents/young adults with attention-deficit/hyperactivity disorder (ADHD). (29) rTMS was administered in 10 sessions over 2 weeks, with 1 week of no TMS between the active and sham phases. The clinical global impression and ADHD-IV scales improved in both conditions over the course of the study, with no significant differences between the active and sham phases.

Bulimia Nervosa

In 2008, Walpoth et al. reported no evidence of efficacy of rTMS in a small trial (n=14) of patients with bulimia nervosa. (30)

Dysphagia

rTMS for the treatment of dysphagia following stroke has been examined in small randomized controlled trials. One study randomized 26 patients to rTMS or sham over the affected esophageal motor area of the cortex. (31) Ten minutes of rTMS over 5 days reduced both dysphagia on the Dysphagic Outcome and Severity scale and disability measured by the Barthel Index. There was a trend for improved hand grip strength in the rTMS group. Blinded assessment showed that the effects were maintained at 1 month and 2 month follow-up. Another study randomized 30 patients with dysphagia following stroke or traumatic brain injury to high-frequency rTMS, low-frequency rTMS, or sham stimulation. (32) Active or sham rTMS was administered bilaterally over the anterolateral scalp over a period of 2 weeks. Swallowing scale scores improved in both the low-frequency and sham groups. Improvement in videofluoroscopic evaluation was greater in the low-frequency rTMS group than the other 2 groups. Blinding of evaluators was not described.

Study in a larger number of subjects is needed to determine the efficacy of this treatment with greater certainty.

Epilepsy

In 2012, Sun et al. reported a randomized double-blind controlled trial of low-frequency rTMS to the epileptogenic zone for refractory partial epilepsy. (33) Sixty patients were randomized into 2 groups; one group received 2 weeks of rTMS at 90% of resting motor threshold and the other group received rTMS at 20% of resting motor threshold. Outcomes were measured for 8 weeks after the end of treatment. With intent-to-treat analysis, high-intensity rTMS resulted in a significant decrease in seizures.
when compared to baseline (from 8.9 per week at baseline to 1.8 per week at follow-up) and when compared to low-intensity rTMS (from 8.6 at baseline to 8.4 per week at follow-up). High-intensity rTMS also decreased interictal discharges (from 75.1 to 33.6 per hour) and improved ratings on the Symptom Checklist-90. These initial results are promising, but require substantiation in additional trials.

Fibromyalgia

A 2012 systematic review included 4 studies on transcranial direct current stimulation and 5 on rTMS for treatment of fibromyalgia pain. (34) Three of the 5 trials were considered to be high quality. Four of the 5 were double-blind randomized controlled trials; the fifth included study was a case series of 4 patients who were blinded to treatment. Quantitative meta-analysis was not conducted due to variability in brain site, stimulation frequency/intensity, total number of sessions, and follow-up intervals, but 4 of the 5 studies on rTMS reported significant decreases in pain. Greater durability of pain reduction was observed with stimulation of the primary motor cortex compared to the dorsolateral prefrontal cortex.

One of the studies included in the systematic review was a small 2011 trial that was conducted in the U.S. by Short et al. (35) Twenty patients with fibromyalgia, defined by the American College of Rheumatology criteria, were randomized to 10 sessions of left prefrontal rTMS or sham TMS along with their standard medications. At 2 weeks after treatment, there was a significant change from baseline in average visual analog scale (VAS) for pain in the rTMS group (from 5.60 to 4.41) but not in the sham-treated group (from 5.34 to 5.37). There was also a significant improvement in depression symptoms in the active group compared to baseline (from 21.8 to 14.10) but not in the sham group (from 17.6 to 16.4). There were no statistically significant differences between the groups in this small trial.

Additional study is needed to determine effective treatment parameters in a larger number of subjects and to evaluate durability of the effect.

Obsessive Compulsive Disorder

Two small (n=18 and 30) randomized sham-controlled trials found no evidence of efficacy for treatment of obsessive compulsive disorder (OCD), although another small sham-controlled trial (n=21) reported promising results with bilateral stimulation of the supplementary motor area. (36-38)
Panic Disorder

In 2013, Mantovani et al. reported a randomized double-blind sham-controlled trial of low-frequency rTMS to the right dorsolateral prefrontal cortex in 21 patients with panic disorder with comorbid major depression. (39) Response was defined as a 40% or greater decrease on the panic disorder severity scale (PDSS) and a 50% or greater decrease on the HAM-D. After 4 weeks of treatment, the response rate for panic was 50% with active rTMS and 8% with sham. There was no significant difference in the response rate for depressive symptoms (25% active rTMS vs. 8% for sham). After an additional 4 weeks of open-label treatment, the response rate was 67% for panic and 50% for depressive symptoms. Five of 12 responders returned for 6-month follow-up and showed sustained improvement.

Parkinson Disease

A systematic review from 2009 included 10 randomized controlled trials with a total of 275 patients with Parkinson disease. (40) Seven of the studies were double-blind, one was not blinded and 2 of the studies did not specify whether the raters were blinded. In studies that used high-frequency rTMS there was a significant improvement on the Unified Parkinson’s Disease Rating Scale (UPDRS) with a moderate effect size of -0.58. For low-frequency rTMS, the results were heterogeneous and did not significantly reduce the UPDRS. The analyzed studies varied in outcomes reported, rTMS protocol, patient selection criteria, demographics, stages of Parkinson disease and duration of follow-up, which ranged from immediate to 16 weeks after treatment.

In 2012, Benninger et al. reported a randomized double-blind sham-controlled trial of brief (6 sec) very-high-frequency (50 Hz) rTMS over the motor cortex in 26 patients with mild to moderate Parkinson disease. (41) Eight sessions of 50 Hz rTMS did not improve gait, bradykinesia, or global and motor scores on the UPDRS compared to the sham-treated group. Activities of daily living were significantly improved a day after the intervention, but the effect was no longer evident at 1 month after treatment. Functional status and self-reported well-being were not affected by the treatment. No adverse effects of the very-high-frequency stimulation were identified.

Another study from 2012 randomized 20 patients with Parkinson disease to 12 brief sessions (6 min) of high-frequency (5-Hz) rTMS or sham rTMS over the leg area of the motor cortex followed by treadmill training. (42) Blinded evaluation showed a significant effect of rTMS combined with treadmill training on
neurophysiological measures, and change in fast walking speed and the timed up and go task. Mean treadmill speed improved to a similar extent in the active and sham rTMS groups.

Additional study with a larger number of subjects and longer follow-up is needed to determine if rTMS improves motor symptoms in patients with Parkinson disease.

Postpartum Depression

Myczkowski et al. conducted a double-blind sham-controlled study of 14 patients with postpartum depression randomized to 20 sessions of active or sham rTMS over the left dorsolateral prefrontal cortex. (43) A positive response to treatment was defined as a reduction of at least 30% in the HAM-D and Edinburgh Postnatal Depression Scale (EPDS). At 2 weeks after the end of treatment, the active rTMS group showed significant improvements in the HAM-D, Global Assessment Scale, Clinical Global Impression and Social Adjustment Scale. The difference in the EPDS (reduction of 39.4% vs. 6.2% for sham) did not reach statistical significance in this small study, and there were marginal cognitive and social improvements. In addition, results were presented as mean values, rather than by the proportion of patients who showed clinically meaningful improvement.

Posttraumatic Stress Disorder

The efficacy of rTMS for posttraumatic stress disorder (PTSD) has been examined in several small randomized controlled trials.

A 2004 study randomized 24 patients with PTSD to 10 sessions of low-frequency (1 Hz), high-frequency (10 Hz) or sham rTMS over the right dorsolateral prefrontal cortex. (44) Blinded assessment 2 weeks after the intervention found that high-frequency rTMS improved the self-reported PTSD checklist (PCL) by 29.3%, the clinician evaluation on the Treatment Outcome PTSD scale by 39.0%, the HAM-D by 25.9%, and the Hamilton Anxiety Rating Scale by 44.1%. Scores for the sham and low-frequency group were not significantly improved.

In 2012, Watts et al reported a double-blind trial with 20 patients randomized to low-frequency rTMS or sham over the right dorsolateral prefrontal cortex. (45) Blinded evaluation at the end of treatment showed clinically significant improvements in the Clinician Administered PTSD Scale (CAPS) and the PCL compared with sham. Depressive and anxiety symptoms also improved in the rTMS group. Six of the 10 rTMS patients showed a degradation of symptoms between the immediate post-treatment assessment and the 2-month post-treatment follow-up.
In another double-blind trial, 30 patients with PTSD were randomized to deep, high-frequency rTMS after brief exposure to a script of the traumatic event, rTMS after a script of a non-traumatic event, or sham stimulation after a brief script of the traumatic event. (46) Patients received 3 treatment sessions per week for 4 weeks, and response was defined as a 50% or greater improvement in CAPS score. Intent-to-treat analysis showed a significant improvement in the total CAPS score in the exposure + stimulation group (24.3) compared to rTMS alone (7.9) or traumatic exposure with sham rTMS (9.1). The greatest improvement was in the intrusive component of the CAPS scale. Heart rate responses to the traumatic script were also reduced over the 4 weeks of treatment. The proportion of patients who showed a response to treatment was not reported and the durability of the response was not assessed.

Conclusions. Several small randomized controlled trials have reported improvement of PTSD with rTMS over the right dorsolateral cortex. Results of high-frequency versus low-frequency stimulation are conflicting, and durability of the response has not been assessed. Additional study is needed.

Schizophrenia

One of the largest areas of TMS research outside of depressive disorders is the treatment of auditory hallucinations in schizophrenia resistant to pharmacotherapy. In 2011, TEC published an Assessment of TMS as an adjunct treatment for schizophrenia. (47) Five meta-analyses were reviewed, along with randomized controlled trials (RCTs) in which measurements were carried out beyond the treatment period. A meta-analysis of the effect of TMS on positive symptoms of schizophrenia (hallucinations, delusions, and disorganized speech and behavior) did not find a significant effect of TMS. Four meta-analyses that looked specifically at auditory hallucinations showed a significant effect of TMS. It was noted that outcomes were evaluated at the end of treatment, and the durability of the effect is unknown. The Assessment concluded that the available evidence is insufficient to demonstrate that TMS is effective in the treatment of schizophrenia.

A 2012 meta-analysis included 17 randomized double-blind sham-controlled trials (n=337) of the effect of rTMS on auditory hallucinations. (48) When measured at the end of treatment, the mean effect size of rTMS directed at the left temporoparietal area was 0.40 (moderate), and the effect size of rTMS directed at all
brain regions was 0.33 (small). For the 5 trials that examined outcomes of rTMS one month after treatment, the effect was no longer significant.

Blumberger et al. examined the efficacy of priming stimulation (6 Hz) prior to low-frequency stimulation (1 Hz) of Heschl’s gyrus within the left temporoparietal cortex. (49) Fifty-four patients with medication-resistant auditory hallucinations were randomized to receive 20 sessions of left-sided stimulation, priming, or sham rTMS. Response rates on the Psychotic Symptoms Rating Scale did not differ between the 3 treatment groups.

A small (n=18) double-blind randomized sham-controlled trial from 2012 found no significant effect of deep rTMS with an H1 coil on auditory hallucinations. (50)

Conclusions: The evidence on rTMS for the treatment of auditory hallucinations in schizophrenia consists of a number of small randomized controlled trials. Evidence to date shows small to moderate effects on hallucinations when measured at the end of treatment, but evidence suggests that the effect is not durable.

**Stroke**

Hsu et al. reported a meta-analysis of the effect of rTMS on upper limb motor function in patients with stroke in 2012. (51) Eighteen randomized controlled trials with a total of 392 patients were included in the meta-analysis. Most of the studies were double blind (n=11) or single blind (n=3). Eight studies applied low frequency (1 Hz) rTMS over the unaffected hemisphere, 5 applied high frequency (5 Hz) rTMS over the affected hemisphere, and 2 used both low- and high-frequency stimulation. Outcomes included kinematic motion analyses (5 trials), hand grip (2 trials), and the Wolf Motor Function Test (2 trials). Meta-analysis of results showed a moderate effect size (0.55) for rTMS on motor outcome, with a greater effect size of rTMS in patients with subcortical stroke (mean effect size, 0.73) compared to non-specified lesion sites (mean effect size, 0.45), and for studies applying low-frequency rTMS (mean effect size, 0.69) compared to high-frequency rTMS (effect size, 0.41). Effect size of 0.5 or greater was considered to be clinically meaningful.

In 2012, Seniow et al. reported a randomized double-blind sham-controlled pilot study of low-frequency rTMS (1 Hz at 90% of resting motor threshold for 30 min) to the contralesional motor cortex combined with physiotherapy in patients with moderate upper extremity hemiparesis following stroke. (52) Power analysis indicated that a sample size of 129 patients would be required to
detect changes in functional motor ability, but only 40 patients met eligibility criteria over the 4 years of the study. Blinded analysis showed no significant difference in hand function or level of neurological deficit between active or sham rTMS when measured either immediately after the 3-week intervention or at 3-month follow-up.

Conclusions. Evidence consists of a number of randomized controlled trials and a meta-analysis of the effect of rTMS on recovery from stroke. Results are conflicting, and efficacy may depend on the location of the stroke and frequency of the rTMS. Additional study is needed to determine whether rTMS facilitates standard physiotherapy in patients with stroke.

**Practice Guidelines and Position Statements**

The Canadian Network for Mood and Anxiety Treatments (CANMAT)

The Canadian Network for Mood and Anxiety Treatments (CANMAT) updated their clinical guidelines on neurostimulation therapies for the management of major depressive disorder in adults. (53) The evidence reviewed supported electroconvulsive therapy (ECT) as a first-line treatment under specific circumstances; when used in patients who have failed to respond to one or more adequate antidepressant medication trials, ECT response rates have been estimated to be 50–60%. The guidelines considered rTMS to be a safe and well-tolerated treatment, with no evidence of cognitive impairment. Based on the 2008 meta-analysis by Lam et al., (5) response (25%) and remission (17%) rates were found to be greater than sham but lower than for other interventions for TRD, leading to a recommendation for rTMS as a second-line treatment. The guidelines indicated that there is a major gap in the evidence base regarding maintenance rTMS, as only one open-label case series was identified.


American Psychiatric Association

In 2010 the American Psychiatric Association published the third edition of The Practice Guideline for the Treatment of Patients With Major Depressive Disorder (55), which states the following.
Evidence for TMS is currently insufficient to support its use in the initial treatment of major depressive disorder.

Based on the results of a multisite randomized sham-controlled clinical trial of high-frequency TMS over the left dorsolateral prefrontal cortex, TMS was cleared by the FDA in 2008 for use in individuals with major depressive disorder who have not had a satisfactory response to at least one antidepressant trial in the current episode of illness. However, another large randomized sham-controlled trial of TMS added to antidepressant pharmacotherapy showed no significant benefit of left dorsolateral prefrontal cortex TMS. In comparisons of actual TMS versus sham TMS, most but not all, recent meta-analyses have found relatively small to moderate benefits of TMS in terms of clinical response. Although the primary studies used in these meta-analyses are highly overlapping and the variability in TMS stimulus parameters and treatment paradigms complicates the interpretation of research findings, these meta-analyses also support the use of high-frequency TMS over the left dorsolateral prefrontal cortex. Lesser degrees of treatment resistance may be associated with a better acute response to TMS.

In comparison with ECT, TMS has been found in randomized studies to be either less effective than ECT or comparable in efficacy to ECT, but in the latter studies TMS was more effective and ECT was less effective than is typically seen in clinical trials. A fewer number of studies have compared cognitive effects of TMS and ECT. One randomized trial found no significant difference between TMS and non-dominant unilateral ECT on performance on neuropsychological tests at 2 and at 4 weeks of treatment, although a small open-label trial reported a greater degree of memory difficulties with ECT than with TMS shortly after the treatment course. Across all studies, TMS was well tolerated and was associated with low rates of treatment dropout. Transient scalp discomfort and headaches were the most commonly reported side effects. In clinical practice, the need for daily TMS could produce logistical barriers for some patients.

The Executive Summary states: “Treatment in the acute phase should be aimed at inducing remission of the major depressive episode and achieving a full return to the patient’s baseline level of functioning [I]. Acute phase treatment may include pharmacotherapy, depression-focused psychotherapy, the combination of medications and psychotherapy, or other somatic therapies such as electroconvulsive therapy (ECT), transcranial magnetic stimulation (TMS), or light therapy...Selection of an initial treatment modality should be influenced by clinical features
(e.g., severity of symptoms, presence of co-occurring disorders or psychosocial stressors) as well as other factors (e.g., patient preference, prior treatment experiences) [I]. Any treatment should be integrated with psychiatric management and any other treatments being provided for other diagnoses [I].” ([I] = Recommended with substantial clinical confidence.)

Agency for Healthcare Research and Quality (AHRQ)

In September, 2011, the Agency for Healthcare Research and Quality (AHRQ) published a Comparative Effectiveness Review of Nonpharmacologic Interventions for Treatment-Resistant Depression in Adults. (56) The Executive Summary contained the following.

Direct evidence. The available head-to-head literature concerning the efficacy of the nonpharmacologic interventions for Tier 1 TRD is limited to two fair trials (both in MDD-only populations). [Tier 1 is studies in which patients specifically had two or more prior treatment failures with medications.] One compared ECT and rTMS, and the other compared ECT and ECT plus rTMS. They showed, with low strength of evidence, no differences between treatment options for depressive severity, response rates, and remission rates.

Indirect evidence. Trials that compared a nonpharmacologic intervention, generally rTMS, VNS, or psychotherapy, with a control or sham procedure in Tier 1 populations were identified. The number of these trials with the same or similar control group was very small, so they could not be pooled quantitatively. They could, however, assess the potential benefits of nonpharmacologic interventions versus controls by calculating mean changes in depressive severity, relative risks of response, and relative risks of remission. rTMS was beneficial relative to controls receiving a sham procedure for all three outcomes (severity of depressive symptoms, response rate, remission rate). rTMS produced a greater decrease in depressive severity (high strength of evidence). Specifically, rTMS averaged a decrease in depressive severity measured by the Hamilton Rating Scale for Depression (HAM-D) of more than 5 points relative to sham control, and this change meets the minimum threshold of the 3-point HAM-D difference that is considered clinically meaningful. Response rates were greater with rTMS than sham (also high strength of evidence); those receiving rTMS were more than three times as likely to achieve a depressive response as patients receiving a sham procedure. Finally, rTMS was also more likely to
produce remission than the control procedure (moderate strength of evidence); patients receiving rTMS were more than six times as likely to achieve remission as those receiving the sham.

Institute for Clinical Systems Improvement

The Institute for Clinical Systems Improvement published a 2012 Health Care Guideline, Major Depression in Adults in Primary Care (57), which states the following.

“While many rTMS studies have been conducted, results are heterogeneous, likely due to small sample sizes and significant variability of anatomical localizations and stimulation intensities and parameters. Compared with early rTMS studies, more recent studies improve upon methodological limitations including active sham treatment mimicking the somatosenory experience of rTMS, masking rTMS administrators and patients to acoustic signals produced by stimulation, and competency certification for outcome evaluators... At this time, a number of treatment and protocol variations for rTMS remain, and the optimum treatment protocol and patient characteristics may not yet be identified. Nonetheless, rTMS is a low-risk and appealing treatment for treatment-refractory depressed patients for whom it is practical and cost-effective.”

Summary

Evidence on repetitive transcranial magnetic stimulation (rTMS) for depression and other psychiatric/neurologic disorders is insufficient to permit conclusions regarding the effect of this technology on health outcomes. The literature on rTMS for treatment-resistant depression is the most developed and includes a number of double-blind randomized sham-controlled short-term trials. Results of these trials show mean improvements of uncertain clinical significance across groups as a whole. The percentage of subjects who show a clinically significant response is reported at approximately 2 to 3 times that of sham controls, with approximately 15% to 25% of patients meeting the definition of clinical response. (26)

The treatment protocols are time intensive, and the patients who are most likely to benefit from treatment are not currently known. Importantly, a number of open issues need to be addressed before this procedure is widely implemented. The available studies do not establish that rTMS is as good as available alternatives, as the vast majority of the trials do not compare rTMS to alternative active treatments. Alternative treatments include a variety of different medication regimens evidence-based
psychotherapies, both of which have demonstrated efficacy. While further research is needed to determine whether the response to acute phase treatment is durable with or without anti-depressant medications, and to provide some information about whether maintenance treatments are needed, and which types of maintenance treatment are most effective, the safety and efficacy of acute phase rTMS for treatment-resistant populations seems to point towards a reasonable and available treatment for an otherwise difficult to treat population. In addition, rTMS has some limited support in professional guidelines for use in TRD.

For other psychiatric/neurologic conditions, the evidence is insufficient to determine whether rTMS leads to improved outcomes. The available clinical trials are small and report mixed results for a variety of conditions other than depression. There are no large, high-quality trials for any of these other conditions.

Coding:

Disclaimer for coding information on Medical Policies

Procedure and diagnosis 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, device or diagnosis 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/HCPCS/ICD-9/ICD-10 Codes

The following codes may be applicable to this Medical policy and may not be all inclusive.

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| HCPCS Codes |
None

**ICD-9 Diagnosis Codes**

Refer to the ICD-9-CM Manual

**ICD-9 Procedure Codes**

N/A

**ICD-10 Diagnosis Codes**

Refer to the ICD-10-CM Manual

**ICD-10 Procedure Codes**

N/A

**Medicare Coverage:**

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.

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

**References:**

1. Blue Cross and Blue Shield Association Technology Evaluation Center (TEC). Transcranial magnetic stimulation for depression. TEC Assessments 2009; Volume 24, Tab 5.

2. Blue Cross and Blue Shield Association Technology Evaluation Center (TEC). Transcranial magnetic stimulation for depression. TEC Assessments 2011; Volume 26, Tab 3.


42. Yang YR, Tseng CY, Chiou SY et al. Combination of rTMS and treadmill training modulates corticomotor inhibition and improves walking in Parkinson disease: a randomized trial. Neurorehabil Neural Repair 2013; 27(1):79-86.


47. Blue Cross and Blue Shield Association Technology Evaluation Center (TEC). Transcranial magnetic stimulation for the treatment of schizophrenia. TEC Assessments 2011; Volume 26, Tab 6.


Policy History:

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http://www.medicalpolicy.hcsc.net/medicalpolicy/activePolicyPage?id(hw77j6nu&corpEn... 7/1/2014
Document updated with literature review. The following changed: 1) rTMS may be considered medically necessary for treatment of major depressive disorder that is resistant to other treatment, when the specific criteria are met; 2) Navigated TMS has been moved to MED205.037 Navigated Transcranial Magnetic Stimulation (nTMS). Title changed from Transcranial Magnetic Stimulation (TMS).

The following was added: Navigated transcranial magnetic stimulation (nTMS) is considered experimental, investigational and unproven. CPT/HCPCS code(s) updated.


New medical document. Transcranial magnetic stimulation (TMS) is considered experimental, investigational and unproven as a treatment of depression and other psychiatric or neurologic disorders including, but not limited to, schizophrenia or migraine headaches. (Coverage is unchanged. This topic was previously addressed on PSY301.000.)

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