

Efficacy and Safety of Transcranial Magnetic Stimulation in the Acute Treatment of Major Depression: A Multisite Randomized Controlled Trial

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Background: We tested whether transcranial magnetic stimulation (TMS) over the left dorsolateral prefrontal cortex (DLPFC) is effective and safe in the acute treatment of major depression.

Methods: In a double-blind, multisite study, 301 medication-free patients with major depression who had not benefited from prior treatment were randomized to active ($n = 155$) or sham TMS ($n = 146$) conditions. Sessions were conducted five times per week with TMS at 10 pulses/sec, 120% of motor threshold, 3000 pulses/session, for 4–6 weeks. Primary outcome was the symptom score change as assessed at week 4 with the Montgomery–Åsberg Depression Rating Scale (MADRS). Secondary outcomes included changes on the 17- and 24-item Hamilton Depression Rating Scale (HAMD) and response and remission rates with the MADRS and HAMD.

Results: Active TMS was significantly superior to sham TMS on the MADRS at week 4 (with a post hoc correction for inequality in symptom severity between groups at baseline), as well as on the HAMD17 and HAMD24 scales at weeks 4 and 6. Response rates were significantly higher with active TMS on all three scales at weeks 4 and 6. Remission rates were approximately twofold higher with active TMS at week 6 and significant on the MADRS and HAMD24 scales (but not the HAMD17 scale). Active TMS was well tolerated with a low dropout rate for adverse events (4.5%) that were generally mild and limited to transient scalp discomfort or pain.

Conclusions: Transcranial magnetic stimulation was effective in treating major depression with minimal side effects reported. It offers clinicians a novel alternative for the treatment of this disorder.

Key Words: Clinical trial, efficacy, major depression, safety, TMS

Major depressive disorder (MDD) is a common, recurrent, and frequently chronic disorder that is a leading contributor to functional impairment and disability (Murray and Lopez 1996). Treatment is often challenging; an estimated 20%–40% of patients do not benefit sufficiently from existing antidepressant interventions including trials of medication and psychotherapy (Greden 2001). A substantial proportion of patients manifest a chronic, treatment-resistant course of illness, resulting in a need for additional treatment options (Rush *et al.*

2006; Trivedi *et al.* 2006). Transcranial magnetic stimulation (TMS) has been proposed as one such alternative (George *et al.* 1997, 1995b; George and Wassermann 1994).

During TMS, a time-varying current is discharged in an insulated coil placed on the scalp surface. This generates a brief dynamic magnetic field that is orthogonal in orientation to current flow in the coil (Amassian *et al.* 1992; Roth *et al.* 1991a, 1991b). The scalp and skull are transparent to the magnetic field, which induces current flow when it reaches a conductive medium such as neural tissue and with it the potential to modulate neural circuitry in a therapeutic fashion.

Several single-center, controlled studies of TMS have been conducted that, in most cases, have supported the hypothesis that TMS manifests antidepressant properties when delivered to the left or right dorsolateral prefrontal cortex (DLPFC; Avery *et al.* 2005; Fitzgerald *et al.* 2003; George *et al.* 1997, 2000; Klein *et al.* 1999; Loo *et al.* 1999, 2001; Pascual-Leone *et al.* 1996). This target was initially selected (George *et al.* 1995b) based on imaging findings implicating this region in the pathophysiology of depression and in antidepressant effects, as well as studies linking specific lesion locations to dysregulation of mood (Bench *et al.* 1992, 1995; Brody *et al.* 2001; George *et al.* 1995a, 1993a, 1993b; Kimbrell *et al.* 2002; Mayberg 2002, 2003; Nobler *et al.* 2001; Sackeim 2001a; Sackeim 2001b; Robinson *et al.* 1984, 1988).

Meta-analyses conducted of this literature have largely concluded that either a slow rate of stimulation (≤ 1 pulse/sec) over the right DLPFC or fast stimulation (at 5–20 pulses/sec) over the left DLPFC have greater antidepressant effects than matched sham stimulation conditions (Burt *et al.* 2002; Couturier 2005; Kozel and George 2002; Martin *et al.* 2002; McNamara *et al.* 2001). Such effects, however, have been questioned in terms of clinical significance, with at least one meta-analysis, using con-

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servative criteria, finding no clear benefit from active TMS (Martin *et al.* 2002). All meta-analyses have acknowledged the limitations of prior work including insufficient intensity and duration of treatment, inferior sham condition, single-center study design, and inadequate sample size.

We conducted a large, multicenter, randomized controlled trial of TMS to address key prior shortcomings. We administered TMS at fixed treatment parameters over the left DLPFC and compared it with a sham TMS intervention that mimicked the active condition to a greater extent than has been the case in previous research. The TMS parameters in the study were selected to achieve the highest feasible dose consistent with current safety guidelines (Wassermann 1998) and patient tolerability. The duration of the TMS course (4–6 weeks) was also longer than in previous studies, providing greater opportunity for cumulative therapeutic and adverse effects to be expressed.

Methods and Materials

Subjects

Eligible subjects were antidepressant medication-free outpatients, aged 18–70, with a DSM-IV diagnosis of MDD, single episode or recurrent, with a current episode duration of 3 years or less. The episode had a Clinical Global Impressions Severity of Illness (CGI-S) score of at least 4 and a total score of at least 20 on the 17-item Hamilton Depression Rating Scale (HAM-D17). Symptom stability was required during a 1-week no-treatment lead-in period, with a HAM-D17 total score of at least 18 and a decrease in score of 25% or less from that observed at the screening assessment. Prior antidepressant treatment during the current episode was assessed using the Antidepressant Treatment History Form (ATHF; Sackeim 2001b). Patients were required to have failed at least one but no more than four adequate antidepressant treatments in this or the most recent episode. Alternatively, patients were eligible if they had marked intolerance to antidepressants as demonstrated by four failed attempts to tolerate an adequate medication trial (lifetime).

Exclusionary criteria for study participation included a lifetime history of psychosis, bipolar disorder, or obsessive-compulsive disorder; posttraumatic stress disorder and eating disorders (if present in the past year); lack of response to an adequate trial of electroconvulsive therapy (ECT); prior treatment with TMS or a vagus nerve stimulator implant; pregnancy; a personal or close family history of a seizure disorder; presence of neurologic disorder or medication therapy known to alter seizure threshold; or presence of ferromagnetic material in or in close proximity to the head. Routine laboratory studies (complete blood count, chemistry, thyroid stimulating hormone), urine toxicology screen, and electrocardiogram were performed at study screening, and subjects were required to be medically stable before entry.

Study Overview

The study was conducted at 23 study sites in the United States (20 sites), Australia (2 sites), and Canada (1 site), with active enrollment extending from January 2004 through August 2005. Institutional review board approval was obtained at all sites. The study was conducted under an Investigational Device Exemption from the U.S. Food and Drug Administration (FDA). All subjects signed an informed consent document before undergoing any study procedures.

The study had three phases: a lead-in phase (1 week, no treatment), a 6-week acute treatment period (daily treatment with TMS or sham), and a taper phase (3 weeks reduced frequency of

TMS or sham, start of antidepressant). Patients were randomized 1:1 to either active TMS or sham TMS. During the acute treatment phase, TMS sessions were scheduled daily in a 5-day sequence, for a maximum of 30 sessions (6 weeks), and typically administered on a Monday through Friday schedule. Then TMS was tapered in a blinded manner in six sessions across 3 weeks during which all patients were titrated onto monotherapy with an antidepressant medication. After 4 weeks of participation in the acute phase, if patients failed to show meaningful clinical benefit (i.e., < 25% reduction in baseline symptoms on the HAM-D17), they could crossover to an open-label, acute treatment extension study. The specific criterion for early study exit was concealed from the investigators' knowledge.

Study Device Description, TMS Session Procedures, and Ratings

The TMS sessions were delivered using the Neuronetics Model 2100 Therapy System investigational device (Neuronetics Inc., Malvern, Pennsylvania). Three separate magnetic coils, similar in weight, external appearance, and acoustic properties when actively pulsed, were used at each site, with one coil unblinded and used as the known active coil to determine motor thresholds. The other two coils differed in that the sham coil had an embedded magnetic shield. The latter limited the magnetic energy reaching the cortex to 10% or less than the active coil but nevertheless allowed the active and sham coils to have similar appearance, placement, and acoustic properties. All treatment personnel were blind as to coil assignment. All efficacy outcome measures were assessed by blinded study personnel (raters) who were not permitted access to the treatment sessions. Raters underwent certification in which their study participation was contingent on demonstrating adequate reliability in the conduct and scoring of interviews to derive HAM-D and MADRS scores. Quality of ongoing ratings was assessed through the use of video monitoring reviewed by an independent expert. Patients were instructed not to disclose any details of the treatment session with the study raters during rating sessions. Furthermore, neither raters nor other study personnel at the specific centers were aware of the primary efficacy measure during the trial. Ratings were administered at baseline and at weeks 2, 4, and 6–9.

Treatment Parameters

Treatment was fixed at 120% magnetic field intensity relative to the patient's observed resting motor threshold (MT), at a repetition rate of 10 magnetic pulses/sec, with a stimulus train duration (on time) of 4 sec and an intertrain interval (off time) of 26 sec. The left DLPFC was the treatment location and was determined by movement of the TMS coil 5 cm anterior to the motor threshold location along a left superior oblique plane with a rotation point about the tip of the patient's nose (George *et al.* 1995b). Spatial coordinates were recorded with a mechanical coil positioning system to ensure placement reproducibility. The MT estimation was repeated weekly by visual observation of thumb or other finger movement (Pridmore *et al.* 1998) using the MT Assist (Neuronetics Inc.). The latter is a standardized, software-based mathematical algorithm that provides an iterated estimate of the MT.

During the first week of the acute phase only, treatment intensity could be adjusted to 110% of MT for tolerability but was then required to return to 120% MT from week 2 onward. A treatment session lasted for 37.5 min for a total of 3000 magnetic pulses delivered per session.

Concomitant Treatments

All patients were free of antidepressants or other psychotropic medications directed at treating depression. Patients were allowed only limited use of either hypnotics or anxiolytics for treatment-emergent insomnia or anxiety, respectively. Up to 14 daily doses (lorazepam 2 mg/day equivalent) only were permitted (of either or both types of medications) during the acute phase.

Efficacy Assessments

The primary efficacy outcome was the difference between active and sham TMS using the last visit MADRS score through week 4 of the acute phase. Secondary outcome measures were the MADRS score at 6 weeks, 24-item and 17-item HAMD scores at 4 and 6 weeks, and categorical endpoints using MADRS, HAMD17, and HAMD24 at 4 and 6 weeks. Response was defined as at least 50% reduction from baseline score. Remission was defined by an absolute scale-specific score, as indicated in Figure 2). Several standardized HAMD factor scores were also derived: including the Depression Core factor (Items 1, 2, 3, 7, 8), the Maier subscale (Items 1, 2, 7, 8, 9, 10), the Gibbons subscale (Items 1, 2, 3, 7, 9, 10, 11, 14), the Anxiety/Somatization factor (Items 10, 11, 12, 13, 15, 17), the Retardation factor (Items 1, 7, 8, 14), and the Sleep factor (Items 4–6). Global clinical status was assessed using the observer rated Clinical Global Impressions Severity of Illness Scale (CGI-S).

Patient-reported outcomes were obtained using the Inventory of Depressive Symptoms—Self Report version (IDS-SR), and the Patient Global Impressions Improvement Scale (PGI-I). Various functional status and quality-of-life measures were collected and will be reported elsewhere.

Safety Assessments

Safety was assessed at every treatment visit by recording spontaneous adverse event reports that were coded using the current version of the *Medical Dictionary for Regulatory Activities*. Additional safety evaluations included targeted assessment of air-conduction auditory threshold at baseline, week 4, and week 6. Cognitive function was assessed periodically and will be the subject of a separate report.

Statistical Methods

Sample size was determined by requiring 90% power and a two-tailed 5% level in detecting a difference between the active and sham conditions with a putative MADRS effect size of approximately .40, based on the standard *t* test method. Efficacy analyses were performed on the strict intent-to-treat sample of all evaluable patients, defined in the protocol as those with a baseline and at least one postbaseline observation available for analysis. The null hypothesis for the primary outcome was tested with an analysis of covariance, using baseline score and ATHF medication resistance level as fixed-effect covariates, adjusting for site differences using a random effect (SAS Institute, Cary, North Carolina). Secondary outcome analyses for continuous measures were conducted in a similar fashion. All analyses were conducted in a last-observation carried forward (LOCF) manner through the indicated time points.

Results

Patient Disposition

Of the 325 patients randomized to TMS or sham, 301 (92.6%) had at least one postbaseline assessment, and their data comprised the a priori-specified analysis set. There were no baseline clinical or demographic differences between the 24 nonevaluable patients and the 301 evaluable subjects. Nonevaluable patients were evenly distributed between the active ($n = 10$) and the sham ($n = 14$) conditions, and there was no systematic difference between these groups in the reasons for discontinuation. Reasons given for study discontinuation through week 4, the primary efficacy time point, are provided in Figure 1.

Through the primary efficacy time point of 4 weeks, the overall discontinuation rate was low and similar in the active TMS (7.7%) and sham TMS (8.2%) treatment groups. Discontinuation because of adverse events of TMS was rare, similar across treatment conditions (4.5% of active TMS vs. 3.4% of sham TMS patients), and most commonly due to scalp discomfort. Beyond the primary efficacy time point, 74 (47.7%) patients in the active

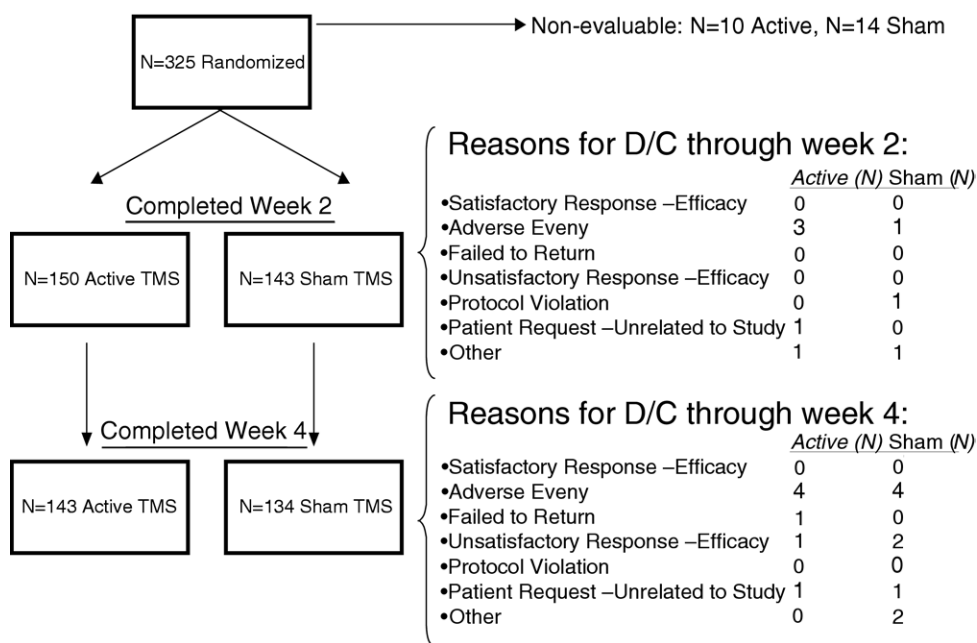


Figure 1. Reasons for study discontinuation through the primary efficacy time point (week 4).

TMS group and 92 (63.0%) patients in the sham TMS treatment group elected to enter the open-label extension study.

Sample Characteristics

Demographic and clinical features were similar in both groups. The CGI-S severity score of 4.7 for the study cohort corresponded to a depression severity between moderately and markedly ill. A minority of the sample reported a current episode duration greater than 2 years. The degree of treatment resistance in the current episode did not differ between groups, with an average of 1.6 failed adequate antidepressant treatments in the current episode and about half the study population failing to benefit from at least two treatments (by ATHF criteria). The degree of functional impairment in the study cohort was substantial, with about half of the sample unemployed and one third on disability because of their mood disorder.

As indicated in Table 1, baseline symptom severity at randomization differed between groups on the primary outcome measure (MADRS, $p = .036$) but not on any of the other symptom scales. This difference arose in the context of the protocol not specifying a minimal symptom severity score on the MADRS for study inclusion, differing in this respect from the HAMD17 (for which a minimum score of 20 was required at the screening visit). Six participants had baseline MADRS scores in the mild

range (< 20 , range 14–19) and were randomized unevenly across the two groups (i.e., 4 to active, and 2 to sham). This resulted in a small but significant difference in baseline symptom scores (i.e., 1.1 point).

Continuous Efficacy Outcome Measures

Efficacy results for continuous outcomes with the MADRS, HAMD24, and HAMD17 are shown in Figure 2.

MADRS

At the primary efficacy time point, week 4, the baseline to endpoint change on the MADRS showed a statistical trend favoring active TMS in the a priori-specified evaluable study population ($p = .057$). Given the observed baseline imbalance on the MADRS, a supplementary analysis was conducted on a study sample that included only those patients ($n = 295$) with a minimum baseline score of 20 on the MADRS, excluding the six patients noted earlier. In this analysis, the baseline to endpoint change was statistically significant at week 4 ($p = .038$). At the secondary efficacy time point, week 6, the baseline to endpoint change on the MADRS continued to show a statistical trend favoring active TMS in the a priori-specified evaluable study population ($p = .057$) and in the subset of patients with a baseline MADRS > 20 ($p = .052$).

Table 1. Demographic Features, Clinical Features, and Baseline and End Point (Weeks 4 and 6) Symptom Scores of the Study Sample

	Active TMS ($n = 155$)	Sham TMS ($n = 146$)	p Value ^a
Demographic Variables			
N (%) female	86 (55.5)	74 (50.7)	.421
Age (years \pm SD)	47.9 \pm 11.0	48.7 \pm 10.6	.509
Ethnic origin, n (%)			
Caucasian	146 (94.2)	131 (89.7)	
Other	9 (5.8)	15 (10.3)	.201
Employment status, n (%)			
Full time	58 (35.6)	45 (28.3)	
Part time	27 (16.6)	31 (19.5)	
Unemployed	78 (47.9)	83 (52.2)	
Receiving disability compensation	28 (32.9)	31 (34.1)	
Clinical Variables			
Recurrent illness course (%)	149 (95.5)	136 (93.8)	.611
Duration of current episode in months, Mean (SD)	13.6 (9.9)	13.2 (9.5)	.728
n (%) of population with current episode > 2 years	36 (23.2)	23 (15.8)	.112
Number of fully adequate antidepressant treatments in current episode	1.6	1.6	.816
Baseline Symptom Scores			
MADRS total score (SD)	32.8 (6.0)	33.9 (5.7)	.036
HAMD17 total score (SD)	22.6 (3.3)	22.9 (3.5)	.508
HAMD24 total score (SD)	30.1 (5.0)	30.5 (4.9)	.568
CGI-Severity (SD)	4.7 (.6)	4.7 (.7)	.594
IDS-SR total score (SD)	42.0 (9.4)	43.4 (9.9)	.197
Week 4 Symptom Scores			
MADRS total score (SD)	27 (11.1)	29.8 (10.1)	.057
HAMD17 total score (SD)	17.4 (6.5)	19.4 (6.5)	.006
HAMD24 total score (SD)	23.4 (8.9)	25.9 (8.8)	.012
Week 6 Symptom Scores			
MADRS total score (SD)	26.8 (12.8)	30 (10.8)	.058
HAMD 17 total score (SD)	17.1 (7.7)	19.6 (7.0)	.005
HAMD 24 total score (SD)	23.2 (10.6)	26 (9.4)	.015

CGI-S, Clinical Global Impressions Severity of Illness; HAMD, Hamilton Depression Rating Scale (17 and 24 item); IDS-SR, Depressive Symptoms—Self Report version; MADRS, Montgomery–Asberg Depression Rating Scale; TMS, transcranial magnetic stimulation.

^a p values represent between group contrasts at baseline except for the weeks 4 and six symptom scores where the p values indicate the contrast of the within group changes from baseline.

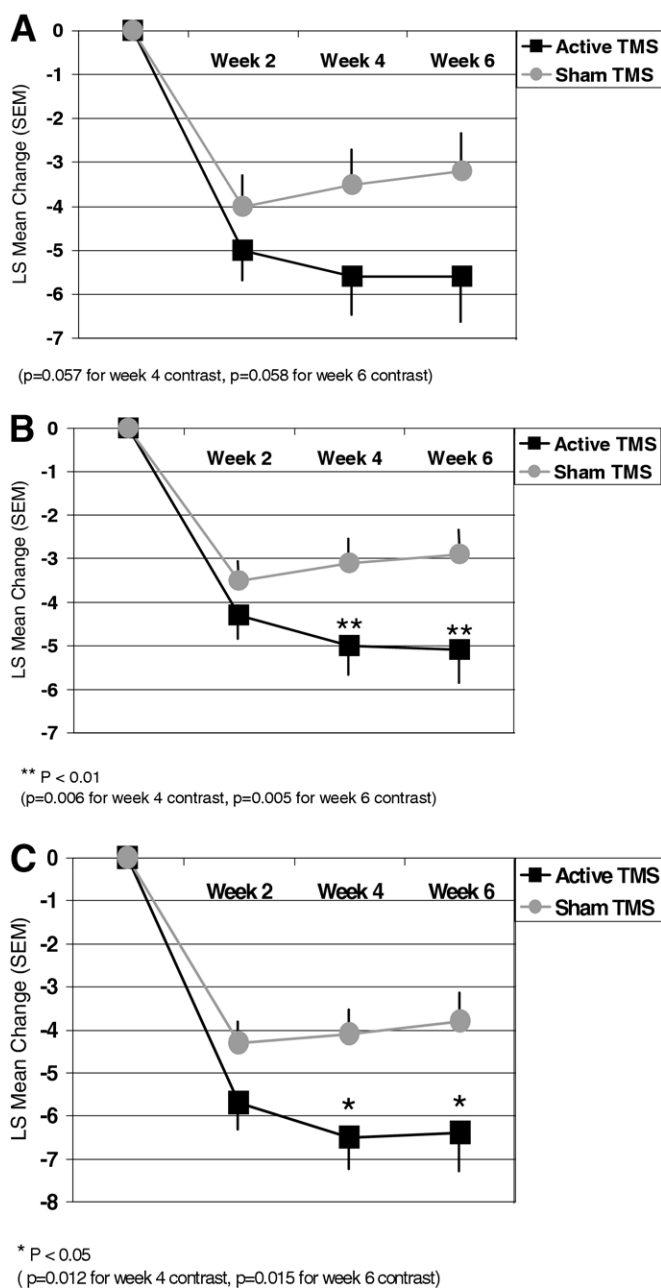


Figure 2. (A) Montgomery–Asberg Depression Rating Scale total score change from baseline during the acute treatment phase. (B) Hamilton Depression Rating Scale (HAMD; 17 item) total score change from baseline during the acute treatment phase. (C) HAMD (24-item) total score change from baseline during the acute treatment phase.

HAMD17 and HAMD24

Results for the HAMD24 and HAMD17 are presented for the a priori–specified evaluable data set only ($n = 301$). No baseline imbalance in symptom severity occurred with these scales, and the findings were unaltered when the six patients with low scores on the MADRS were excluded (data available from the authors on request). At the primary efficacy time point, week 4, the baseline to endpoint change on both the HAMD17 and the HAMD24 yielded a significant main effect of treatment group favoring active TMS ($p = .006$ and $p = .012$, respectively). This outcome was sustained at the secondary efficacy time point,

week 6, with a significant advantage in favor of active TMS ($p = .005$ for HAMD17, $p = .015$ for HAMD24). As seen in Figure 2A–C, it is notable that the time course of clinical effect showed a sustained improvement across the acute treatment phase for the active TMS group, whereas the sham TMS group showed a pattern of early change that dissipates at the later time points.

CGI-S

Clinician-rated global illness severity showed greater improvement with active TMS compared with sham TMS as early as week 2 of the acute treatment phase ($p = .047$) and continued through the primary efficacy time point of week 4 ($p = .009$) and the secondary efficacy time point of week 6 ($p = .012$).

Categorical Outcomes

Response Rates. As seen in Figure 3A–C, at the week 4 primary efficacy time point, the response rate as defined by the three symptom-rating scales (MADRS, HAMD17, and HAMD24) was higher with active compared with sham TMS. This effect was sustained through the secondary efficacy time point at week 6.

Remission Rates. As seen in Figure 3A–C, at the week 4 primary efficacy time point, a significant difference in remission rates was not detected. At the secondary efficacy time point of week 6, however, remission rate was higher with active compared with sham TMS on the MADRS and HAMD24.

HAMD Factor Scores. The pattern of change in the HAMD17 scale was examined further by the assessment of previously established factor scores for the HAMD. These results are shown in Table 2 and demonstrate significantly better outcomes for active TMS compared with sham TMS on core depression symptoms (Core Depression Factor, Maier Factor, and Gibbons Factor), anxiety symptoms (Anxiety/Somatization Factor), and vegetative symptoms (Retardation Factor) at weeks 4 and 6.

Patient-Reported Mood Symptoms and Global Improvement. Patient-reported mood change and global improvement were assessed using the IDS-SR and the PGI–Improvement scales. The pattern of change followed the clinician-reported measures but showed a less robust effect. The IDS-SR exhibited a trend toward improved outcome in active compared with sham TMS at the primary efficacy time point, week 4 ($p = .058$), which was maintained at the secondary efficacy time point, week 6 ($p = .053$). There was no statistically significant separation between the groups on the PGI-I at either time point.

Antidepressant Effects During TMS Taper. During this phase, response and remission rates of the active TMS group improved incrementally. With the MADRS, the response rate increased from 23.9% to 27.7% and the remission rate from 14.2% to 20.6% for. In contrast, addition of medication to the sham TMS group produced little meaningful clinical change during this phase: a change from 12.3% to 13.7% for the response rate and from 5.5% to 8.9% for remission on the MADRS.

Safety Outcomes

Spontaneous Adverse Events. There was a higher incidence of scalp discomfort and pain with active than sham TMS (Table 3). These events were generally reported as mild or moderate in severity and diminished rapidly in incidence after the first treatment week. Scalp discomfort had the potential to compromise the study blind, and a separate analysis was conducted to examine the relationship between clinical outcome and the experience of cutaneous discomfort. The findings were negative regarding an association between any of these adverse event terms and the primary outcome measure (data not shown).

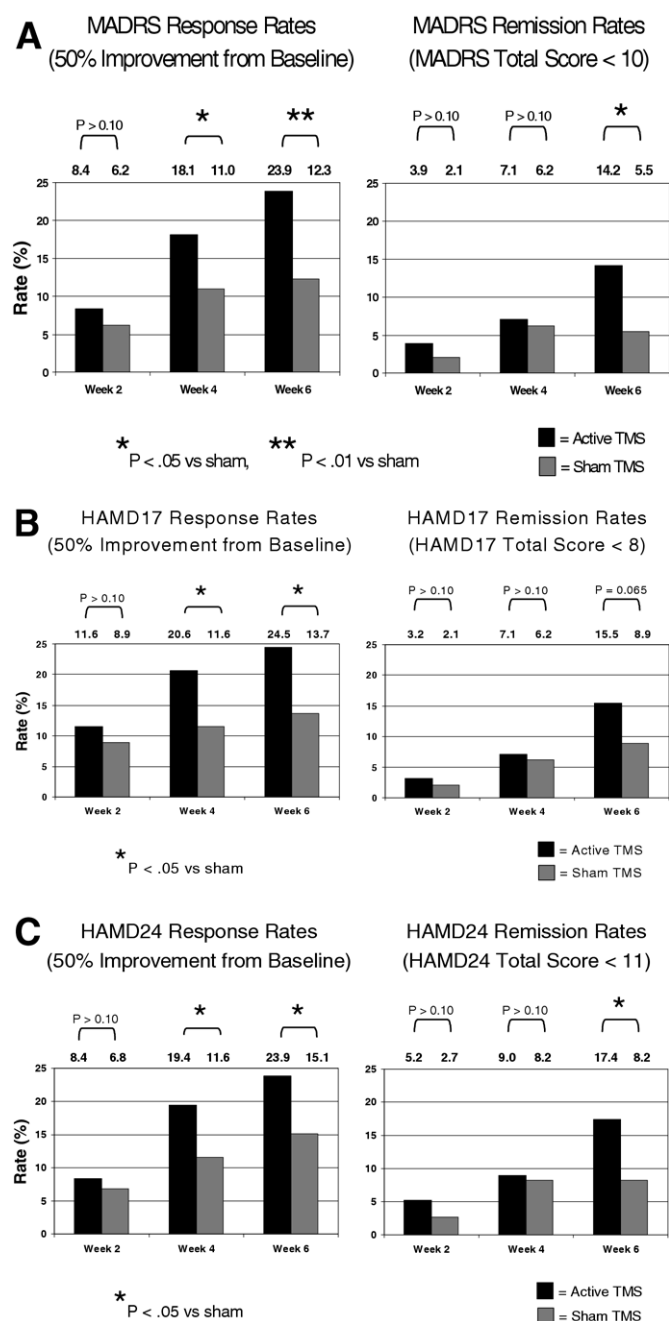


Figure 3. (A) Montgomery–Asberg Depression Rating Scale categorical outcome assessments during the acute treatment phase. (B) Hamilton Depression Rating Scale (HAMD; 17 item) categorical outcome assessments during the acute treatment phase. (C) HAMD (24 item) categorical outcome assessments during the acute treatment phase.

Serious Adverse Events. There were no deaths in this study, and no seizures were reported. During the acute treatment phase, 16 serious adverse events were reported, 9 in the active TMS group and 7 in the sham TMS group. Events reflecting disease-related exacerbation were the most common serious adverse events. These included suicidality (1.9% with sham vs. .6% with active TMS), exacerbation of depression (1.9% with sham vs. .6% with active), and a single suspected suicide gesture (in the sham group). Overall risk of exacerbation of suicidality was evaluated by determining the proportion of patients in either

group who increased in score on the suicide item of the HAMD (item 3) from a value of 0 or 1 at baseline to a value of 3 or 4 at any time point during the acute treatment phase. Cumulatively, 10 events meeting this criterion were observed in the sham TMS group compared with 1 event in the active TMS group.

Audiometry. All subjects used earplugs during the treatment sessions. No differences in air-conduction thresholds were detected between or within treatment groups across the acute treatment phase of the study (data not shown, to be presented in a subsequent report).

Discussion

This is the first large, multisite, randomized controlled trial of daily left prefrontal TMS in medication-free patients with major depression who had failed to receive adequate benefit from prior antidepressant treatment. The findings indicate that TMS, administered at these parameters for a period of 4–6 weeks, is safe and effective in the treatment of major depression.

The pattern of symptom improvement was consistent across the various study outcome measures. Active treatment with TMS was significantly superior to sham TMS treatment for the change in mean symptom score using the HAMD17 and HAMD24 at weeks 4 ($p = .006$, $p = .012$ respectively) and 6 ($p = .005$, $p = .015$). After correction for the baseline score imbalance, the MADRS also showed this pattern ($p = .038$). Clinically important change, as reflected in terms of the categorical outcomes of response and remission, was also achieved in a substantial portion of patients. At 6 weeks, the active TMS group was about twice as likely to have achieved remission compared with the sham TMS group (MADRS:14.2% vs. 5.2%, HAM-D17: 15.5% vs. 7.1%, HAMD24; 17.4% vs. 8.2%). Improvement on the self-report IDS scale was at the level of a trend ($p = .058$ week 4; $p = .053$ week 6). Clinical outcomes were observed in the setting of a favorable tolerability profile, with less than 5% of patients on active TMS discontinuing treatment because of adverse effects by the primary efficacy endpoint of 4 weeks.

The trajectory of improvement, as indicated in Figure 2, implies that more than 2 weeks of TMS, compared with sham, is required in this population before a significant improvement is detected. Similarly, it appears that an additional 2 weeks of TMS beyond the initial 4 weeks, as indicated in Figure 3, can have an important clinical impact. The remission rates doubled during that period of time.

Table 2. HAMD Factor Scores—Contrasts Between Active TMS and Sham TMS during the Acute Treatment Phase

HAMD Rating Scale Factor Score	Week 2	Week 4	Week 6
Depressive Symptoms			
Core depression factor	>.100	.012	.008
Maier factor	>.100	.003	.003
Gibbons factor	>.100	.007	.006
Anxiety Symptoms			
Anxiety/somatization factor	>.100	.025	.023
Vegetative symptoms			
Retardation factor	.057	.007	.003
Sleep factor	>.100	>.100	>.100

HAMD, Hamilton Depression Rating Scale; TMS, transcranial magnetic stimulation. All contrasts, except for the sleep factor, reflect a superior outcome at weeks 4 and 6 for the active TMS compared with the sham TMS treatment condition. *p* values shown for analysis of covariance model; post hoc contrast between active and sham TMS at the time points indicated.

Table 3. Adverse Events Occurring in the Active Treatment Group at a Rate of 5% or More and at Least Twice the Rate for Sham (with ME-Coded Preferred Terms Shown)

Body System Preferred term	Active TMS (n = 165) n (%)	Sham TMS (n = 158) n (%)
Eye disorders		
Eye pain	10 (6.1)	3 (1.9)
Gastrointestinal Disorders Toothache	12 (7.3)	1 (.6)
General Disorders and Site Administration Conditions		
Application site discomfort	18 (10.9)	2 (1.3)
Application site pain	59 (35.8)	6 (3.8)
Facial pain	11 (6.7)	5 (3.2)
Musculoskeletal and connective tissue disorders		
Muscle twitching	34 (20.6)	5 (3.2)
Skin and subcutaneous tissue disorders		
Pain of skin	14 (8.5)	1 (.6)

MedDRA, *Medical Dictionary for Regulatory Activities*.

In this study, TMS was well tolerated and safe. The dropout rate for any reason from active TMS was 7.7% at 4 weeks, and discontinuation specifically because of side effects was 4.5%. This is lower than the discontinuation rates generally reported from clinical trials of marketed antidepressants. Adverse events reported were principally limited to scalp discomfort or pain within the confines of the TMS session itself and were mostly transient phenomena in the first weeks of the TMS course. In fact, the incidence of headache did not differ between active and sham TMS conditions.

This study also suggests that the upper limit on the safe administration of TMS may be somewhat greater than we currently suspect. Despite TMS being administered here at 120% of motor threshold and 3000 pulses/session, an elevated rate of serious adverse events relative to sham was not detected. The most serious side effect that has been reported with TMS is a seizure (Anderson *et al.* 2006), and none were observed in this study. No adverse effects on mood in terms of either treatment-emergent suicidality or treatment-emergent mania or hypomania were observed. In fact, the frequency of treatment-emergent suicidal ideation was numerically lower in the active TMS group (1 event observed with active vs. 10 events observed with sham during the acute treatment phase).

How can the clinical significance of these results be placed in context for the practitioner? In answering this question, it is important to consider the treatment resistance of the population of patients studied. On average, the patients included had an average of 1.6 failed adequate trials in the current episode of major depression, and nearly half had failed to benefit from 2 or more treatments. The importance of prior resistance to antidepressant treatment is to diminish significantly the likelihood of responding to subsequent interventions. This pattern has been demonstrated with respect to outcomes with electroconvulsive therapy (Prudic *et al.* 1990, 1996; Sackeim *et al.* 1990), pharmacotherapy (Trivedi *et al.* 2006), and vagus nerve stimulation (Sackeim *et al.* 2001).

The most comprehensive demonstration of this in the recent literature is from the large, open-label, seminaturlistic STAR*D trial (Trivedi *et al.* 2006). In that program, there was a progressive reduction in remission rates with each stage of resistance. In fact, patients who had failed two prior antidepressant treatments experienced a mean remission rate (defined as a HAMD17 score < 8) in level 3 treatment of only 16.2% after a 10- to 12-week

course of treatment (Fava *et al.* 2006). In the controlled trial results reported here, at the end of 6 weeks of treatment with TMS alone, patients experienced a remission rate of 15.5%, and this increased to 22.6%, by the same HAMD17 criteria, after 9 weeks during the taper phase of the study. Given that controlled trials generally report somewhat lower clinical response and remission rates than are seen in open-label experience, the results reported here compare favorably to those seen in similarly treatment-resistant patients in the STAR*D reports.

Controlled trials, where an active antidepressant medication is measured against its relevant within-study control, provide the most appropriate comparison for the TMS results. In the large analysis of the FDA database of approved antidepressant medications, Khan and colleagues (2000) noted that the mean percentage reduction from baseline in total HAMD17 score across the entire data set of antidepressants was 40.7% for active treatment and 30.9% for placebo. On average, this represents an overall relative advantage of about 10% in the reduction of total score from baseline on the HAMD17 when comparing active antidepressant to placebo treatment. By comparison, in this study, at 4 weeks, active TMS treatment resulted in a reduction in HAMD17 score of approximately 23% compared with 15% in the sham group. This represents an overall relative advantage of about 8% for active TMS compared with sham in reduction of the total HAMD17 score. Similarly, application of the metric of number needed to treat (NNT; Kraemer and Kupfer 2006) as an indicator of effect size, yielded an NNT for TMS of 11 at week 4 and 9 at week 6. Thase *et al.* (2005), recently reported a pooled estimate of response rates in a large sample ($N = 1795$) of patients treated with bupropion, various selective serotonin reuptake inhibitors, or placebo and noted an overall response rate in active-treated patients of 62.8% compared with 50.8% in placebo treated patients, yielding an NNT in that sample of eight. Thus, by these metrics, the antidepressant efficacy of TMS is comparable to that of standard pharmacotherapy.

This study has several limitations that should be considered when interpreting the results. Although a rigorous method was used to assess prior treatment resistance, namely, the ATHF, this method relies on retrospective report. Therefore, greater certainty regarding the level of treatment resistance would have been obtained from a prospective, open-label antidepressant lead-in phase and subsequently enrolling only nonresponders to that intervention. This prospective method, however, requires considerably more time and cost to implement and is therefore not commonly used. The data documented in this trial support a moderate level of treatment resistance for the study population. Moreover, the observed response and remission rates with the sham TMS intervention were very low, providing additional evidence of the treatment resistance of this patient population.

The study employed an innovative approach to sham methodology that represents a clear advance over prior work. Special efforts were made to match the active and sham conditions in procedure, sound, and sensation as closely as possible, while substantially limiting the exposure of the cortex to the actual magnetic field, as discussed in the study methods. Formal query of patients and treaters to assess the adequacy of the blind, however, was not conducted. Results from prior studies indicate that previously TMS-naïve patients primarily base judgment of whether they received an active or sham TMS procedure on clinical outcome (Fitzgerald *et al.* 2003). A secondary analysis (results to be presented in a subsequent report) also indicated that scalp discomfort with active TMS did not correlate with

treatment outcome. Thus, unblinding of the active condition is an unlikely explanation for the therapeutic advantage of active TMS.

The primary efficacy endpoint in this study was at week 4 of the 6-week acute treatment phase. At or after that time point, if clinically justified, patients were eligible to enter the open-label extension study. Complete randomization was thus only truly present through the primary efficacy time point. Therefore, outcomes at time points after week 4 should be interpreted in this context. On the other hand, because the treatment assignment blind was maintained throughout the study, it is worth noting that active TMS subjects continuing through the end of the acute phase and into the taper phase of the study showed persistent and perhaps accumulating benefit in comparison to their continuing counterparts in the sham TMS group. Finally, the method of coil positioning used to identify the left DLPFC was not based on guidance by means of an magnetic resonance imaging–assisted neuronavigational method. Rather, we used a probabilistic surface anatomy approach targeting 5 cm anterior to the motor threshold location, like most clinical trials of TMS. This approach may not fully account for individual differences in brain anatomy (Herwig *et al.* 2001, 2003).

In conclusion, TMS administered over the left DLPFC using the parameters reported here for a period of up to 6 weeks was effective in treating major depression and with a good tolerability profile. These results indicate that TMS offers clinicians a novel alternative in the treatment of this disorder (Sackeim 2001a).

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- Amassian VE, Eberle L, Maccabee PJ, Cracco RQ (1992): Modelling magnetic coil excitation of human cerebral cortex with a peripheral nerve immersed in a brain-shaped volume conductor: The significance of fiber bending in excitation. *Electroencephalo Clin Neuro* 85:291–301.
- Anderson B, Mishory A, Nahas Z, Borckardt JJ, Yamanaka K, Rastogi K, George MS (2006): Tolerability and safety of high daily doses of repetitive transcranial magnetic stimulation in healthy young men. *J ECT* 22:49–53.
- Avery DH, Holtzheimer PE 3rd, Fawaz W, Russo J, Neumaier J, Dunner DL, *et al.* (2005): A controlled study of repetitive transcranial magnetic stimulation in medication-resistant depression. *Biol Psychiatry*, September 1 [Epub ahead of publication].
- Bench CJ, Frackowiak RS, Dolan RJ (1995): Changes in regional cerebral blood flow on recovery from depression. *Psychol Med* 25:247–261.
- Bench CJ, Friston KJ, Brown RG, Scott LC, Frackowiak RS, Dolan RJ (1992): The anatomy of melancholia—focal abnormalities of cerebral blood flow in major depression. *Psychol Med* 22:607–615.
- Brody AL, Saxena S, Stoessel P, Gillies LA, Fairbanks LA, Alborzian S, *et al.* (2001): Regional brain metabolic changes in patients with major depression treated with either paroxetine or interpersonal therapy: preliminary findings [see comment]. *Arch Gen Psychiatry* 58:631–640.

- Burt T, Lisanby SH, Sackeim HA (2002): Neuropsychiatric applications of transcranial magnetic stimulation: A meta analysis. *Int J Neuropsychopharmacol* 5:73–103.
- Couturier JL (2005): Efficacy of rapid-rate repetitive transcranial magnetic stimulation in the treatment of depression: A systematic review and meta-analysis [see comment]. *J Psychiatry Neurosci* 30:83–90.
- Fava M, Rush AJ, Wisniewski SR, Nierenberg AA, Alpert JE, McGrath PJ, *et al.* (2006): A comparison of mirtazapine and nortriptyline following two consecutive failed medication treatments for depressed outpatients: A STAR*D report. *Am J Psychiatry* 163:1161–1172.
- Fitzgerald PB, Brown TL, Marston NA, Daskalakis ZJ, De Castella A, Kulkarni J (2003): Transcranial magnetic stimulation in the treatment of depression: A double-blind, placebo-controlled trial. *Arch Gen Psychiatry* 60:1002–1008.
- George MS, Ketter TA, Parekh PI, Horwitz B, Herscovitch P, Post RM (1995a): Brain activity during transient sadness and happiness in healthy women. *Am J Psychiatry* 152:341–351.
- George MS, Ketter TA, Post RM (1993a): SPECT and PET imaging in mood disorders. *J Clin Psychiatry* 54:6–13.
- George MS, Ketter TA, Post RM (1993b): SPECT and PET imaging in mood disorders. *J Clin Psychiatry* 54(suppl):6–13.
- George MS, Nahas Z, Molloy M, Speer AM, Oliver NC, Li XB, *et al.* (2000): A controlled trial of daily left prefrontal cortex TMS for treating depression [see comment]. *Biol Psychiatry* 48:962–970.
- George MS, Wassermann EM (1994): Rapid-rate transcranial magnetic stimulation and ECT. *Convuls Ther* 10:251–254; discussion 255–258.
- George MS, Wassermann EM, Kimbrell TA, Little JT, Williams WE, Danielson AL, *et al.* (1997): Mood improvement following daily left prefrontal repetitive transcranial magnetic stimulation in patients with depression: A placebo-controlled crossover trial. *Am J Psychiatry* 154:1752–1756.
- George MS, Wassermann EM, Williams WA, Callahan A, Ketter TA, Bassar P, *et al.* (1995b): Daily repetitive transcranial magnetic stimulation (rTMS) improves mood in depression. *Neuroreport* 6:1853–1856.
- Greden JF (2001): The burden of recurrent depression: Causes, consequences, and future prospects. *J Clin Psychiatry* 22:5–9.
- Herwig U, Satrapi P, Schonfeldt-Lecuona C (2003): Using the international 10-20 EEG system for positioning of transcranial magnetic stimulation. *Brain Topogr* 16:95–99.
- Herwig U, Schonfeldt-Lecuona C, Wunderlich AP, von Tiesenhäusen C, Thielscher A, Walter H, Spitzer M (2001): The navigation of transcranial magnetic stimulation. *Psychiatry Res* 108:123–131.
- Khan A, Warner HA, Brown WA (2000): Symptom reduction and suicide risk in patients treated with placebo in antidepressant clinical trials: an analysis of the Food and Drug Administration database [see comment]. *Arch Gen Psychiatry* 57:311–317.
- Kimbrell TA, Ketter TA, George MS, Little JT, Benson BE, Willis MW, *et al.* (2002): Regional cerebral glucose utilization in patients with a range of severities of unipolar depression. *Biol Psychiatry* 51:237–252.
- Klein E, Kreinin I, Chistyakov A, Koren D, Mecz L, Marmur S, *et al.* (1999): Therapeutic efficacy of right prefrontal slow repetitive transcranial magnetic stimulation in major depression: A double-blind controlled study [see comment]. *Arch Gen Psychiatry* 56:315–320.
- Kozel A, George M (2002): Meta-analysis of left prefrontal repetitive transcranial magnetic stimulation (rTMS) to treat depression. *J Psychiatr Pract* 8:270–275.
- Kraemer HC, Kupfer DJ (2006): Size of treatment effects and their importance to clinical research and practice. *Biological Psychiatry* 59:990–996.
- Loo C, Mitchell P, Sachdev P, McDermott B, Parker G, Gandevia S (1999): Double-blind controlled investigation of transcranial magnetic stimulation for the treatment of resistant major depression. *Am J Psychiatry* 156:946–948.
- Loo C, Sachdev P, Elsayed H, McDermott B, Mitchell P, Wilkinson M, *et al.* (2001): Effects of a 2- to 4-week course of repetitive transcranial magnetic stimulation (rTMS) on neuropsychologic functioning, electroencephalogram, and auditory threshold in depressed patients. *Biol Psychiatry* 49:615–623.
- Martin JL, Barbanj MJ, Schlaepfer TE, Clos S, Perez V, Kulisevsky J, Gironell A (2002): Transcranial magnetic stimulation for treating depression. *Cochrane Database of Systematic Reviews* 2: CD003493.
- Mayberg HS (2002): Modulating limbic-cortical circuits in depression: Targets of antidepressant treatments. *Semin Clin Neuropsychiatry* 7:255–268.
- Mayberg HS (2003): Positron emission tomography imaging in depression: a neural systems perspective. *Neuroimag Clin North Am* 13:805–815.
- McNamara B, Ray JL, Arthurs OJ, Boniface S (2001): Transcranial magnetic stimulation for depression and other psychiatric disorders [see comment]. *Psychol Med* 31:1141–1146.
- Murray C, Lopez A (1996): *The Global Burden of Disease: A Comprehensive Assessment of Mortality and Disability from Diseases, Injuries and Risk Factors in 1990 and Projected to 2020*. Cambridge, MA: Harcourt University Press.
- Nobler MS, Oquendo MA, Kegeles LS, *et al.* (2001): Decreased regional brain metabolism after ECT. *Am J Psychiatry* 158:305–308.
- Pascual-Leone A, Rubio B, Pallardo F, Catala MD (1996): Rapid-rate transcranial magnetic stimulation of left dorsolateral prefrontal cortex in drug-resistant depression [see comment]. *Lancet* 348:233–237.
- Pridmore S, Fernandes Filho JA, Nahas Z, Liberatos C, George MS (1998): Motor threshold in transcranial magnetic stimulation: A comparison of a neurophysiological method and a visualization of movement method. *J ECT* 14:25–27.
- Prudic J, Haskett RF, Mulsant B, Malone KM, Pettinati HM, Stephens S, *et al.* (1996): Resistance to antidepressant medications and short-term clinical response to ECT [see comment]. *Am J Psychiatry* 153:985–992.
- Prudic J, Sackeim HA, Devanand DP (1990): Medication resistance and clinical response to electroconvulsive therapy. *Psychiatry Res* 31:287–296.
- Robinson RG, Boston JD, Starkstein SE, Price TR (1988): Comparison of mania and depression after brain injury: Causal factors. *Am J Psychiatry* 145:172–178.
- Robinson RG, Kubos KL, Starr LB, Rao K, Price TR (1984): Mood disorders in stroke patients: Importance of location of lesion. *Brain* 107:81–93.
- Roth BJ, Cohen LG, Hallett M (1991a): The electric field induced during magnetic stimulation. In Lacey WJ, Cracco RQ, Barker AT, Rothwell J, editors. *Magnetic Motor Stimulation: Basic Principles and Clinical Experience* (EEG Suppl. 43). Amsterdam: Elsevier Science, 268–278.
- Roth BJ, Saypol JM, Hallett M, Cohen LG (1991b): A theoretical calculation of the electric field induced in the cortex during magnetic stimulation. *Electroencephalogr Clin Neurophysiol* 81:47–56.
- Rush AJ, Trivedi MH, Wisniewski SR, Stewart JW, Nierenberg AA, Thase ME, *et al.* (2006): Bupropion-SR, sertraline, or venlafaxine-XR after failure of SSRIs for depression. *New Engl J Med* 354:1231–1242.
- Sackeim HA (2001a): Functional brain circuits in major depression and remission [comment]. *Arch Gen Psychiatry* 58:649–650.
- Sackeim HA (2001b): The definition and meaning of treatment-resistant depression. *J Clin Psychiatry* 16:10–17.
- Sackeim HA, Prudic J, Devanand DP, Decina P, Kerr B, Malitz S (1990): The impact of medication resistance and continuation pharmacotherapy on relapse following response to electroconvulsive therapy in major depression. *J Clin Psychopharmacol* 10:96–104.
- Sackeim HA, Rush AJ, George MS, Marangell LB, Husain MM, Nahas Z, *et al.* (2001): Vagus nerve stimulation (VNS) for treatment-resistant depression: Efficacy, side effects, and predictors of outcome. *Neuropsychopharmacology* 25:713–728.
- Thase ME, Haight BR, Richard N, Rockett CB, Mitton M, Modell JG, *et al.* (2005): Remission rates following antidepressant therapy with bupropion or selective serotonin reuptake inhibitors: a meta-analysis of original data from 7 randomized controlled trials. *J Clin Psychiatry* 66:974–981.
- Trivedi MH, Rush AJ, Wisniewski SR, Nierenberg AA, Warden D, Ritz L, *et al.* (2006): Evaluation of outcomes with citalopram for depression using measurement-based care in STAR*D: Implications for clinical practice. *Am J Psychiatry* 163:28–40.
- Wassermann EM (1998): Risk and safety of repetitive transcranial magnetic stimulation: Report and suggested guidelines from the International Workshop on the Safety of Repetitive Transcranial Magnetic Stimulation, June 5–7, 1996. *Electroencephalogr Clin Neurophysiol* 108:1–16.