**An ERP Study of Multidimensional Source Retrieval in Depression**

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**Abstract**

Major Depressive Disorder (MDD) affects episodic memory, but few functional imaging studies have examined the underlying neural mechanisms. Therefore, we collected event-related potentials (ERPs) from 24 unmedicated adults with MDD and 24 healthy controls during source memory retrieval. At encoding, neutral words were presented on the left or right (perceptual source) in the context of animacy or mobility judgments (conceptual source). Mobility judgments were made more slowly than animacy judgments, consistent with prolonged semantic analysis. At retrieval, accuracy for words from the mobility task was characterized by a *Group* x *Cue* interaction, with the MDD group showing a benefit for conceptual vs. perceptual retrieval not seen in the controls. Similarly, left parietal ERPs elicited by words from the mobility task were characterized by *Group* x *Cue* interactions from 400-800 ms and 800-1400 ms post-stimulus. In these intervals, the MDD group showed higher amplitude ERPs for conceptual vs. perceptual retrieval, but the controls did not. In the MDD group, the magnitude of these ERP effects was negatively correlated with self-reported sleep disruption. By contrast, accuracy for words from the animacy task was characterized by a main effect of *Cue*:conceptual source accuracy was worse than perceptual source accuracy across both groups, and ERP difference waves tracking this effect revealed sustained negative potentials over fronto-central scalp. These data highlight the sensitivity of conceptual (but not perceptual) source memory to the extent of semantic processing at encoding, and indicate that source memory in MDD covaries with left parietal ERPs linked to recollection.

Keywords: source memory, retrieval, ERP, recollection, depression

**1. Introduction**

Major Depressive Disorder (MDD) is known for its association with biased emotional memory. Relative to healthy controls, depressed adults typically show excellent memory for negative stimuli and poor memory for positive stimuli (Burt et al., 1995; Dillon et al., 2014; Hamilton and Gotlib, 2008; Matt et al., 1992), and we have proposed that the positive memory deficit may reflect disruption of brain dopamine systems (Dillon, 2015). Importantly, though, one need not use emotional stimuli to detect an episodic memory deficit in depression. In a meta-analysis of 22 neuropsychological studies, Zakzanis and colleagues (1998) found strong negative effects of depression on episodic memory tests conducted with neutral stimuli. Similarly, Airaksinen et al. (2004) used neutral words to demonstrate impaired free and cued recall in 187 depressed adults recruited from the community. Finally, a meta-analysis of over 30 years of work with the Cambridge Neuropsychological Test Automated Battery (CANTAB) indicated that, relative to healthy adults, depressed individuals show significantly impaired episodic memory, despite the fact that the CANTAB tasks use neutral stimuli (Rock et al., 2014).

What underlies the memory impairment for neutral stimuli in depression? Poor encoding is an obvious candidate. Along these lines, Airaksinen et al. (2004) found that although memory accuracy was consistently lower in depressed vs. healthy adults, both groups showed a similar benefit for cued vs. free recall, implying that the depressed group was able to use cues to enhance retrieval but simply had stored less material in the first place. A more nuanced result was presented by Zakzanis et al. (1998), who found that the effect of depression on memory varied by encoding task: larger negative effects were seen for tasks that provided less structure during learning (e.g., memorization of uncategorized vs. categorized word lists). The cognitive initiative framework offers a principled account of such results (Hertel, 1997; Hertel and Hardin, 1990). The framework’s core hypothesis is that depressed individuals can control attention and use strategies to enhance encoding but that—in the absence of external support or emotionally compelling material—they often fail to do so. To test this hypothesis, Hertel and Rude (1991) presented neutral words and sentence frames to depressed and healthy participants, who were asked to judge whether each word fit its frame. Participants were assigned to either focused or unfocused conditions. In the focused condition, each word was flashed briefly and then replaced by its frame for 8 s, at which point the participant restated the word and indicated whether or not there was a quality fit. In the unfocused condition, the words and frames were presented concurrently (so that the participants did not have to store the words in working memory but could view them alongside the frames), the participants were not required to restate the words, and responses could be made at any time. In short, the unfocused condition was designed to permit mind-wandering and rumination, both of which are common in depression (e.g., Nolen-Hoeksema, 1991). Because mind-wandering and rumination should impair encoding, Hertel and Rude predicted a negative effect of depression specifically in the unfocused condition. Indeed, free recall was characterized by a *Group* x *Task* interaction, with depressed adults remembering fewer words than controls from the unfocused task but not the focused task. Thus, depression impaired memory when encoding was unconstrained, but the provision of a task that engaged attention and encouraged elaborative encoding reduced that deficit.

The cognitive initiative framework also applies to retrieval. Depressed adults typically show larger deficits for recall than for recognition (Burt et al., 1995), and when recognition memory is analyzed to estimate contributions made by recollection vs. familiarity (or controlled vs. automatic processing), depression invariably impairs the former more than the latter (e.g., Hertel and Milan, 1994; MacQueen et al., 2002). The cognitive initiative framework explains these data by pointing to the greater need for controlled attention, effortful searching, and post-retrieval monitoring during free recall vs. recognition, and in support of recollection vs. familiarity. To test this account, Hertel and Milan (1994) conducted studies in which healthy and depressed participants completed successive implicit (homophone spelling) and explicit (recognition) memory tests for the same words. When the experimenters did not draw attention to the fact that identical words appeared on both tests, healthy participants were more likely than depressed adults to notice this regularity and exploit it to improve recognition (i.e., by referring back to the implicit test). However, once the relationship between the tests was highlighted, the group difference in recognition memory disappeared. Moreover, when participants were given a strategy for improving their memory, depressed adults showed an especially large benefit. These results provide additional support for the claim that depression impairs episodic memory for neutral stimuli when the encoding and/or retrieval tasks are relatively unstructured, but that engaging tasks (or detailed instructions concerning helpful strategies) can rescue performance.

Given the extensive and elegant behavioral work on these issues, the paucity of relevant neuroscientific data is surprising. In particular, although there are many studies of hippocampal volumes in depression (for review, see MacQueen and Frodl, 2011) and some functional imaging investigations of encoding (e.g., Bremner et al., 2004; Dillon et al., 2014; Dillon and Pizzagalli, 2013; Hamilton and Gotlib, 2008), there are remarkably few studies on the neuroscience of memory retrieval in MDD. This does not reflect lack of desire—about a decade ago, the National Institutes of Mental Health, Aging, and Neurological Disorders and Stroke called for integrated neuroscientific research on depression and memory (Steffens et al., 2006). However, despite dozens of event-related potential (ERP) and functional magnetic resonance imaging (fMRI) studies of episodic retrieval in healthy adults (Eichenbaum et al., 2007; Rugg and Curran, 2007; Rugg and Vilberg, 2013), no similar literature has emerged in MDD.

The current study was designed to address this gap. Because depression affects recollection more than familiarity—and given the difficulties associated with imaging free recall—we elected to conduct an ERP investigation of source memory in MDD. Source memory refers to conscious retrieval of the spatiotemporal details that define an encoding episode (Johnson et al., 1993). Importantly, source memory depends heavily on recollection—although familiarity can play an important role (Mollison and Curran, 2012)—and there is evidence that it is disrupted in depression (Degl’Innocenti and Bäckman, 1999). We used a design that recruits neural systems engaged during conceptual and perceptual source retrieval (Bergström et al., 2013; Dobbins and Wagner, 2005; Simons et al., 2005a). At study, participants viewed neutral words presented on the left or right above a question specifying an animacy or mobility judgment. At test, they were cued to recall the presentation side (perceptual source, “Side” cue) and encoding task (conceptual source, “Question” cue). A recent fMRI/ERP study in healthy adults (Bergström et al., 2013) found that both conceptual and perceptual retrieval elicited the most well-studied ERP marker of recollection: a positive deflection over parietal scalp that extends from about 400-800 ms post-stimulus, typically with a left hemisphere maximum (Rugg and Curran, 2007). Both forms of retrieval also activated the precuneus and elicited a negative polarity ERP maximal over posterior electrodes that is referred to as the late posterior negativity, or LPN (Cycowicz et al., 2001; Johansson and Mecklinger, 2003; Mecklinger et al., 2007). Intriguingly, the LPN extended over left frontal scalp during conceptual retrieval, and this was mirrored by fMRI activation in left dorsolateral PFC. Related fMRI studies confirmed that left and medial PFC regions were more strongly activated during conceptual vs. perceptual source retrieval (Simons et al., 2005a,b).

Based on this prior work, we expected depression to impair conceptual source memory, for two reasons. First, MDD is associated with volumetric losses in medial PFC (Treadway et al., 2015), and to the extent that medial PFC is especially important for conceptual vs. perceptual retrieval (Simons et al., 2005a,b), we reasoned that conceptual retrieval would be especially disrupted in MDD. Second, rumination is negatively associated with episodic memory in depression (Hertel, 1998), and rumination activates PFC circuits particularly active during conceptual retrieval (Cooney et al., 2010); thus, we expected heightened rumination in MDD to preferentially disrupt conceptual retrieval. Because prior work has presented the retrieval data collapsed over different encoding tasks (e.g., Bergström et al., 2013; Simons et al., 2005a), we did not expect conceptual or perceptual retrieval to vary with the encoding tasks in either group.

However, it quickly became apparent that this expectation was in error. As described below, although both encoding tasks required analysis of the semantic properties of the words, the mobility task was more difficult and elicited longer response times (RTs), which had consequences for retrieval. First, and consistent with the cognitive initiative framework, we found that when sustained analysis at encoding was followed by conceptual retrieval—i.e., when words from the mobility task were presented under the Question cue—performance in the MDD group was particularly good. Second, we found that conceptual retrieval was strongly shaped by the encoding tasks but perceptual retrieval was not, and this was true in both groups. As detailed below, these two behavioral results were associated with markedly different ERP effects. Thus, this experiment provides insight into the neural activity mediating source memory in MDD when encoding and retrieval are well-supported, as well as evidence that the ERPs elicited during successful conceptual (vs. perceptual) retrieval vary according to the cognitive processes that were engaged at encoding.

**2. Materials and Methods**

**2.1. Participants and self-report**

Participants (18-62 years old, right-handed, no neurological or unstable medical conditions) were recruited from the community and compensated $25/hour, following a protocol approved by the Partners HealthCare Human Research Committee. They were screened by phone or online, at which time the Beck Depression Inventory II (BDI; Beck et al., 1996) was administered. Individuals were invited to participate in the MDD group provided they endorsed symptoms consistent with a current Major Depressive Episode, had a BDI-II score ≥ 14 (the accepted cut-off for mild depression; Beck et al., 1996), and reported no other Axis I psychopathology with the exception of generalized anxiety, social anxiety, and/or specific phobia, all of which are highly comorbid with MDD. Controls had to report no current or past Axis I psychopathology. On the day of the ERP experiment, we assessed psychiatric status with the MINI International Neuropsychiatric Interview, version 6.0 (Sheehan et al., 1998). Depressed adults had to again report current depression, no history of other DSM-IV Axis I diagnosis (except generalized anxiety, social anxiety, or specific phobia), and no medication use in the past two weeks (six weeks for fluoxetine, six months for neuroleptics). Thirty-four controls and 26 depressed adults completed the ERP session. Data from 10 controls and 2 depressed adults were excluded due to excessive artifacts (see below), leaving 24 individuals per group.

Following the EEG session, we administered the BDI-II again, along with the Mood and Anxiety Symptom Questionnaire (MASQ; Watson et al., 1995), the Ruminative Responses Scale (RRS; Treynor et al., 2003), and the Pittsburgh Sleep Quality Index (PSQI; Buysse et al., 1989). These probe symptoms of depression and anxiety, trait rumination, and sleep quality over the last month, respectively. The BDI, MASQ, and RRS are commonly used to assess cognitive and affective aspects of depression, with the MASQ also providing insight into anxiety. The PSQI was included because sleep disruption affects neurocognitive processes relevant to episodic retrieval, including executive function in general and the activation of parietal regions implicated in episodic retrieval in particular (Chee et al., 2006; Durmer and Dinges, 2005; McEwen, 2006). Moreover, there is substantial evidence of sleep disruption in depression and other psychiatric disorders (e.g., Deldin et al., 2006; Tsuno et al., 2005; Wulff et al., 2010). Thus, we expected negative relationships between these measures on one hand and both memory accuracy and ERP indices of successful retrieval on the other. Finally, the Wechsler Test of Adult Reading (WTAR; Holdnack, 2001) was used to estimate IQ. One control did not complete the MASQ and one depressed participant did not complete the PSQI. WTAR data from four non-native English speakers (2 controls, 2 MDD) were not analyzed, as results from the WTAR may be invalid in this population.

**2.2. Task**

The task was programmed in PsychoPy (Peirce, 2009). Due to a hardware change, RT data were not recorded for one control and one depressed participant.

**2.2.1. Stimuli.** We used the MRC Psycholinguistic Database (Coltheart, 1981) to select 25 words from four categories: “living/immobile” (e.g., *oak*), “non-living/immobile” (e.g., *shed*), “living/mobile” (e.g., *dog*), and “non-living/mobile” (e.g., *kite*). ANOVA yielded no significant differences for number of letters (mean±S.D.; 5.27±1.29) or syllables (1.52±0.50), frequency (35.58±79.02), concreteness (598.87±20.18), or imageability (596.80±25.31), *ps* > 0.064. Words are listed in the Supplement.

**2.2.2. Encoding.** The task included six encoding-retrieval cycles. Each encoding block included 16 trials (Figure 1, *left*) in which a word appeared on the left or right above one of two questions: “living/non-living?” (animacy task) or “mobile/immobile?” (mobility task). Participants responded by pressing a button. A jittered interval (500-2000 ms) separated the trials.

**2.2.3. Counting.** Immediately after encoding, a 3-digit number (e.g., 931) was shown and participants counted backwards from that number in steps of three for 30 s. Counting served to disrupt rehearsal and clear working memory (Reitman et al., 1974).

[PLEASE INSERT FIGURE 1 ABOUT HERE]

**2.2.4. Retrieval.** Each block comprised 48 trials that included a cue, a word, and a response screen (Figure 1, *right*). On 16 trials each, the cue was “Side” or “Question” and the word came from the preceding encoding block; these trials probed perceptual and conceptual source retrieval, respectively. On the remaining 16 trials the cue was “Odd/Even”, the word was a numeral between “one” and “ninety-six”, and the participant judged parity. All trials involved reading a cue, interpreting it, and retrieving information, but on Odd/Even trials retrieval was directed at semantic rather than episodic memory. Thus, comparing ERP data from Side or Question trials to Odd/Even trials should isolate activity mediating episodic retrieval. Presentation order of words and cues was random. The response screen consisted of ‘RESPOND’ printed above the word with the numbers 1-5 printed below and corresponding to a choice and level of confidence (Figure 1, *right*). As in other studies of multidimensional source memory (Starns and Hicks, 2005), “guess” was included as a response option. Participants were asked to select this when they could not recover any information favoring one source over the other, so that analyses focused on hits should not be contaminated with guesses. A jittered interval (500-2000 ms) separated the trials.

**2.3. EEG Recording**

The EEG was recorded during retrieval with a 128-sensor HydroCel GSN Electrical Geodesics Inc (EGI) net (sample rate: 1000 Hz, 0.02–100 Hz). Data were referenced to vertex and impedances were kept below 45 kΩ when possible (maximum: 75 kΩ).

**2.4. Behavioral Analysis**

The behavioral data were cleaned by dropping trials with no response or where *log*(RT) exceeded the participant’s mean±3SD (< 4% of encoding trials, < 1% of retrieval trials). Analysis involved *t*-tests andmixed-model Type III ANOVAs implemented in the R software (R Core Team, 2015) library *afex* (Singmann et al., 2016). For both encoding and retrieval, accuracy was computed as percent correct. For RT, ANOVAs were computed on *log* transformed data because of improved fit to normality, but the figures and descriptive statistics use untransformed RT data for interpretability.At encoding, *Group* x *Task* (mobility, animacy) x *Side* (left, right) ANOVAs were run for accuracy and correct RT. At retrieval, between-groups *t­*-tests were first used to compare accuracy, confidence, and correct RT on Odd/Even trials, to verify that MDD did not affect performance in this control condition. Next, a *Group* x *Cue* x *Task* ANOVA was run on the number of guesses in each condition. Finally, responses on Question and Side trials were analyzed by running *Group* x *Cue* x *Task* ANOVAs for accuracy, confidence, and correct RT. Alpha was set at 0.05.

**2.5. ERP Analysis**

**2.5.1. Pre-processing.** Pre-processing was conducted with EEGLAB (Delorme and Makeig, 2004) and ERPLAB (Lopez-Calderon and Luck, 2014) toolboxes for MATLAB (MathWorks, Natick). EEG data were merged, re-referenced to the average of all electrodes, and filtered (0.1-30 Hz). Bad channels were interpolated, independent components analysis was used to remove activity reflecting blinks, HEOG, and EKG, and the cleaned data were time-locked to word onsets and segmented (-200 to 2000 ms). The pre-stimulus interval was used for baseline correction, and segments where any raw value or the maximum-minimum voltage difference (200 ms intervals, 100 ms sliding window) exceeded 100 μV were rejected. We used *a priori* criteria of > 18 bad channels or more than 50% of trials rejected (Luck, 2014) to exclude excessively noisy datasets (10 controls, 2 MDD). The mean number of clean segments in each bin defined by *Group* x *Cue* x *Task* ranged from 21-28 for source hits.Guesses were excluded and there were too few clean segments for analyzing misses. Thus, the analysis was focused on correct responses, a common approach in this literature (Bergström et al., 2013; Dobbins and Wagner, 2005; Han et al., 2012; Simons et al., 2005a).

**2.5.2. Group-level analyses**. We conducted two main ERP analyses. In the first, we computed “Question minus Odd/Even” and “Side minus Odd/Even” difference waves, collapsed over the encoding tasks for each participant, and then compared the MDD and control groups. This approach has been used previously (Bergström et al., 2013, Simons et al., 2005a,b), and it allowed us to test our *a priori* prediction that depression would selectively affect conceptual retrieval. The second analysis was intended to parallel the source accuracy results, which revealed a *Group* x *Cue* interaction for words from the mobility task but a main effect of *Cue* for the animacy task. To isolate cue effects while holding encoding processes constant, we computed “Question minus Side” difference waves separately for words from the mobility and animacy tasks in each participant, and then we compared the groups at each level of task.

All within and between-group analyses were statistically analyzed by submitting the difference waves to mass univariate analysis (Groppe et al., 2011a,b) and focusing on mean amplitudes from 400-800 ms, 800-1400 ms, and 1400-2000 ms. The 400-800 ms interval was selected to capture left parietal ERP effects consistently associated with recollection (Rugg and Curran, 2007), with the latter two windows intended to encompass the LPN seen during source recollection (Bergström et al., 2013; Johansson and Mecklinger, 2003). Mass univariate analysis is widely used in fMRI research (Friston et al., 1995) and here entailed one-sample (within-group analysis) or two-sample (between-group analysis) *t*-tests at each electrode. By examining every electrode over multiple time windows, this approach makes better use of the spatiotemporal richness of ERP data than traditional methods. To correct for multiple comparisons, we used cluster-based permutation (Groppe et al., 2011a). All electrodes within 4 cm of each other were considered neighbors, and neighboring electrodes significant at *p* < 0.05 (uncorrected) were considered clusters. The use of a 4 cm inter-electrode distance is appropriate for detecting neighboring electrodes in regions where they are relatively widely spaced (e.g., over parietal scalp) on the 128 channel EGI net (Song et al., 2015). The sum of all *p*-values in a cluster constituted its mass. We then performed 2500 between-participant permutations of the data, each time randomly assigning participants to the two groups without replacement, computing the group difference, and then selecting the most extreme cluster mass to generate a distribution of observed cluster sizes under the null hypothesis of no group difference (Bullmore et al., 1999) (for within-group analyses, the permutation was made between conditions). This null distribution was used to estimate the probability of observing the clusters actually detected when the control and MDD groups were compared. Only clusters significant at *p* < 0.05 (corrected) are reported.

**2.6. Individual Differences**

Across the groups, we used Pearson correlations to examine relationships between source accuracy, confidence, and left parietal ERPs associated with recollection. Within the depressed group, we examined relationships between source accuracy, confidence, left parietal ERPs, sleep quality, the severity of depressive and anxious symptoms, and brooding rumination.

**3. Results**

**3.1. Demographics**

There were no group differences in gender, age, education, or estimated IQ (Table 1). Relative to controls, the MDD group endorsed poorer sleep, more rumination, and more symptoms of depression and anxiety, with the mean BDI-II score indicating moderate depression.

[PLEASE INSERT TABLE 1 ABOUT HERE]

**3.2. Behavior**

**3.2.1. Encoding**. The encoding data are presented in Figure 2. For response accuracy (Figure 2A), the only significant result was a main effect of *Task*, *F*(1, 46) = 13.03, *p* < 0.001, reflecting lower accuracy for mobility vs. animacy judgments. Neither the main effect of *Group* nor any interaction involving this factor approached significance, *F*s < 1. For correct RT (Figure 2B), the main effect of *Task* was again significant, *F*(1, 44) = 50.20, *p* < 0.001, with slower responses for mobility vs. animacy judgments. The RT analysis also revealed a *Group* x *Task* x *Side* interaction, *F*(1, 44) = 8.02, *p* = 0.007, but separate ANOVAs for words presented on the left and right did not reveal significant *Group* x *Task* interactions, *F*s < 3.76, *p*s > 0.05. Examining Figure 2B, the 3-way interaction appears to reflect the fact that while RTs were consistently numerically shorter in the MDD group, this difference was more pronounced for the animacy task for words shown on the left, and for the mobility task for words shown on the right. In summary, the mobility task was more difficult than the animacy task, as judged by lower encoding accuracy and slower RTs, but there were no reliable group differences.

[PLEASE INSERT FIGURE 2 ABOUT HERE]

**3.2.2. Retrieval: Odd/Even trials**. There were also no group differences on Odd/Even trials, which were characterized by extremely high accuracy (controls: 98.44±3.92%; MDD: 99.13±1.37%; *t*(46) = -0.82, *p* = 0.42), fast correct RTs (controls: 867.72±301.94 ms; MDD: 781.28±227.83 ms; *t*(44) = 1.10, *p* = 0.28), and highly confident replies (percentage of high confidence trials: controls: 99.61±0.68%; MDD: 99.87±0.35%; *t*(46) = -1.68, *p* = 0.10).

**3.2.3. Retrieval: guessing**. Five participants (three controls, two MDD) never guessed and a further 12 participants (8 controls, 4 MDD) did not guess in at least one cell of the design. Data from the remaining participants (13 controls, 18 MDD) are shown in Figure 3 and were included in the *Group* x *Cue* x *Task* ANOVA; because only one guess was recorded on an Odd/Even trial, that condition was omitted. The ANOVA yielded an effect of *Cue*, *F*(1, 29) = 7.75, *p* = 0.009, reflecting fewer guesses under the Question vs. Side cue, and an effect of *Task*, *F*(1, 29) = 15.98, *p* < 0.001, reflecting fewer guesses following the mobility vs. the animacy task.

[PLEASE INSERT FIGURE 3 ABOUT HERE]

Neither the main effect of *Group* nor any interactions involving *Group* were significant, but there was a trend for a *Group* x *Cue* x *Task* interaction, *F*(1, 29) = 3.62, *p* = 0.07. Inspection of Figure 3 suggested a pronounced cue effect for words from the mobility task in the MDD group. Indeed, a *t*-test confirmed that, for words from the mobility task, depressed participants guessed less frequently under the Question vs. the Side cue, *t*(17) = -3.29, *p* = 0.004, *d* = 0.77; the cue effect was not reliable for words from the animacy task, *t*(17) < 1. In controls there was no reliable cue effect for either encoding task considered alone (*t*s < 1.33, *p*s > 0.20). Moreover, in the MDD group, words from the mobility task presented under the Question cue elicited fewer guesses than words from the animacy task shown under the Question cue, *t*(17) = -3.74, *p* = 0.002, *d* = 0.82, or words from the animacy task shown under the Side cue, *t*(17) = -3.59, *p* = 0.002, *d* = 0.85. In sum, all participants guessed less to words from the mobility vs. animacy task and in response to the Question vs. Side cue, and in the MDD group, words from the mobility task presented under the Question cue elicited especially few guesses.

**3.2.4. Retrieval: source accuracy**. The source accuracy results are given in Figure 4A; the left and right panels show data for words from the mobility and animacy tasks, respectively. There were main effects of *Cue*, *F*(1, 46) = 20.33, *p* < 0.001, and *Task*, *F*(1, 46) = 30.43, *p* < 0.001, which were qualified by two interactions: *Cue* x *Task*, *F*(1, 46) = 21.69, *p* < 0.001, and *Group* x *Cue*, *F*(1, 46) = 5.49, *p* = 0.02. The *Cue* x *Task* interaction reflected the fact that, when the data were collapsed across the groups, accuracy was lower under the Question vs. Side cue for words from the animacy task, *t*(47) = -6.08, *p* < 0.001, but not for words from the mobility task, *t*(47) = 0.25, *p* = 0.80. An alternative way to view this interaction is to note that accuracy under the Question cue was significantly better for words from the mobility task vs. the animacy task, t(47) = 5.56, *p* < 0.001, while accuracy under the Side cue did not differ significantly by encoding task, *t*(47) = 1.89, *p* = 0.064.

The cause of the *Group* x *Cue* interaction was less clear as there were no significant group differences for accuracy under the Question or Side cue when the data were collapsed across the encoding tasks, *t*s < 1.16, *p*s > 0.25. Inspection of Figure 4A suggested that the cue effect varied by group for words from the mobility task, but not the animacy task. Consistent with this impression—and despite the lack of a three-way interaction (*F* < 1)—there was a significant *Group* x *Cue* effect for words from the mobility task, *F*(1, 46) = 5.92, *p* = 0.02. The interaction was not significant for words from the animacy task, *F* < 1, where accuracy was characterized by a strong main effect of *Cue*, *F*(1, 46) = 36.81, *p* < 0.001.

The nature of these results is highlighted in Figure 4B, which plots “Question minus Side” accuracy difference scores. For words from the mobility task (left panel), the difference score was marginally greater than zero in the MDD group, *t*(23) = 1.77, *p* = 0.090, but marginally lower than zero in the controls, *t*(23) = -1.67, *p* = 0.108, and a between-groups *t*-test was significant, *t*(46) = 2.43, *p* = 0.019, *d*  = 0.70. By contrast, for words from the animacy task (right panel), the difference scores were more negative than zero in both groups, *t*s < -4.29, *p*s < 0.001, and there was no group difference, *t*(46) = 0.92, *p* = 0.36, *d* = 0.27. For both groups, Question minus Side difference scores were more positive for words from the mobility vs. the animacy task, *t*s > 2.7, *ps* < 0.02.

[PLEASE INSERT FIGURE 4 ABOUT HERE]

In summary, the cue effect varied by group for words from the mobility task: depressed adults showed better accuracy under the Question vs. Side cue, but the controls did not. By contrast, the cue effect was stable across groups for words from the animacy task, where accuracy was significantly lower under the Question vs. Side cue.

**3.2.5. Retrieval: source confidence**. Figure 5A shows the confidence data. These were characterized by a main effect of *Task*, *F*(1, 46) = 7.91, *p* = 0.007, reflecting higher confidence in response to words from the mobility vs. animacy task, as well as a main effect of *Cue*, *F*(1, 46) = 24.85, *p* < 0.001, that was qualified by a *Group* x *Cue* interaction, *F*(1, 46) = 4.24, *p* = 0.05. The interaction emerged because there was no group difference in the percentage of high confidence responses under the Question cue, *t*(46) *<* 1, but depressed adults were less confident than controls under the Side cue, *t*(46) = 2.25, *p* = 0.03, *d* = 0.65. Exploratory *t*-tests confirmed that, in the MDD group, confidence was higher under Question vs. Side for words from the mobility task, *t*(23) = 5.35, *p* < 0.001, *d* = 1.10, and the animacy task, *t*(23) = 4.36, *p* < 0.001, *d* = 1.01. Thus, in contrast to the accuracy and guessing data, the cue effect on confidence did not vary by encoding task in the MDD group.

[PLEASE INSERT FIGURE 5 ABOUT HERE]

**3.2.6. Retrieval: RT**. RT data are shown in Figure 5B. The only significant effect was a main effect of *Cue*, with slower responses to Question vs. Side, *F*(1, 44) = 267.92, *p* < 0.001. *Group* was not significant, *F* <1, nor were any interactions with *Group*, *F*s < 1.64, *p*s > 0.20.

**3.2.7. Overall behavioral summary**. At encoding, mobility judgments were made more slowly and less accurately than animacy judgments. This appears to have influenced retrieval as both groups guessed less and responded more confidently to words from the mobility vs. the animacy task. Furthermore, participants responded more slowly, guessed less, and were more confident under the Question vs. Side cue. For source accuracy, the cue effects varied by encoding task. Both groups were less accurate under the Question cue vs. the Side cue in response to words from the animacy task (i.e., main effect of *Cue*). By contrast, accuracy in response to words from the mobility task was characterized by a *Group* x *Cue* interaction: in depressed adults, but not controls, accuracy was better under the Question vs. Side cue. Moreover, depressed participants guessed least frequently in response to words from the mobility task presented under the Question cue. Thus, the mobility task and Question cue led to few guesses and confident responding in all participants, and the combination of these factors boosted accuracy and sharply reduced guessing in the MDD group.

**3.3. ERPs**

**3.3.1. Conceptual and perceptual retrieval, collapsed over encoding task**.To test our *a priori* hypothesis, we conducted between-group tests on “Question minus Odd/Even” and “Side minus Odd/Even” difference waves, collapsed across the encoding tasks as in prior studies. We expected group differences in the former but not the latter contrast, but in fact we found no group differences at all (smallest cluster *p* = 0.29). Thus, we present the results collapsed across groups in Figures 6 and 7.

[PLEASE INSERT FIGURE 6 AND TABLE 2 ABOUT HERE]

Figure 6 depicts the “Question minus Odd/Even” contrast and Table 2 lists electrodes where condition effects were observed. As shown in the top panel, from 400-800 ms there were two clusters of differential activity. As expected, Question hits elicited more positive ERPs than Odd/Even hits over left parietal electrodes, consistent with many prior studies of recollection (Rugg and Curran, 2007). In addition, there was a relative negativity for Question hits vs. Odd/Even hits over right frontal electrodes in this time window. As shown in the middle and bottom panels, later intervals (800-1400 ms and 1400-2000 ms) were characterized by sustained negative polarity potentials for Question hits relative to Odd/Even hits over left frontal and right occipital sites. The left frontal result echoes prior work linking this potential specifically to conceptual retrieval (Bergström et al., 2013), while the LPN over posterior sites is consistently seen in many studies of both conceptual and perceptual source memory (Bergström et al., 2013; Johansson and Mecklinger, 2003; Mecklinger et al., 2007).

For the “Side minus Odd/Even” contrast (Figure 7, Table 3), no reliable differences between conditions were seen from 400-800 ms, but strong effects were observed from 800-1400 ms and 1400-2000 ms. In these windows, a sustained LPN in response to Side hits was evident over the posterior midline, extending from anterior parietal to occipital sites. Robust activity over the posterior midline during retrieval of spatial information is consistent with both prior findings from this task (Bergström et al., 2013) and the larger “PMAT” framework (Ritchey et al., 2015), which describes reliable activation of posterior midline (PM) vs. anterior temporal (AT) structures during retrieval of contextual information in general and spatial information in particular. Thus, both *a priori* contrasts yielded sensible results in line with prior findings, but neither revealed any group differences.

[PLEASE INSERT FIGURE 7 AND TABLE 3 ABOUT HERE]

**3.3.2. *Question* minus *Side*, animacy task**. Our second ERP analysis was designed to more closely track the source accuracy results. Accuracy for words from the animacy task was worse under the Question vs. Side cue in both groups (Figure 4A, right panel). To probe the neural correlates of this effect, we computed “Question minus Side” difference waves for words from the animacy task. A between-groups test revealed no reliable differences (smallest cluster *p* = 0.20), thus we present data collapsed across groups in Figure 8 (also see Table 4). There was a broadly distributed negativity that was focused over left fronto-central scalp from 400-800 ms, dispersed over bilateral fronto-central scalp from 800-1400 ms, and separated into left fronto-central and right centro-parietal clusters from 1400-2000 ms. Inspection of waveforms revealed a stable pattern: relative to Side hits, Question hits elicited more negative potentials, with below-baseline activity especially evident from 1400-2000 ms.

[PLEASE INSERT FIGURE 8 AND TABLE 4 ABOUT HERE]

**3.3.3. *Question* minus *Side*, mobility task**. Finally, we computed “Question minus Side” difference scores for words for the mobility task and compared responses across the two groups. As shown in Figure 9 (see also Table 5), this contrast was associated with group differences over left centro-parietal scalp between 400-800 ms (Figure 9, top) and 800-1400 ms (Figure 9, middle). In these intervals, the depressed and healthy groups generated similar responses for Question hits, but the depressed group showed a weaker response for Side hits. Indeed, follow-up *Group* x *Cue* ANOVAs on mean amplitudes averaged over all electrodes in these clusters (Figure 9, bottom) revealed *Group* x *Cue* interactions in both intervals (*F*s > 14, *p*s < 0.001). These reflected significant control > MDD effects for Side hits (*t*s > 2.2, *p*s < 0.025, *d*s > 0.67), but not Question hits (*t*s < 0.7, *p*s > 0.52, *d*s <0.19). Moreover, the MDD group generated higher amplitude ERPs for Question vs. Side hits in both time windows (*t*s > 3.1, *p*s < 0.006, *d*s > 0.54), but controls showed the opposite pattern—larger ERPs for Side vs. Question hits (*t*s > 2.0, *ps* < 0.056, *ds* >0.20). This ERP pattern is similar to the pattern seen for accuracy (Figure 4).

[PLEASE INSERT FIGURE 9 AND TABLE 5 ABOUT HERE]

**3.4 Individual Differences**

To look for brain/behavior relationships across the groups, we first computed the mean amplitude of the left centro-parietal “Question minus Side” ERP difference waves for words from the mobility task from 400-800 ms and 800-1400 ms, averaging over the electrodes that showed group differences in the top and middle panels of Figure 9, respectively. The amplitudes of the ERPs in these intervals were highly correlated, *r* = 0.84, *p* < 0.001. These values were also positively correlated with “Question minus Side” accuracy and confidence difference scores, although the relationships were modest (**400-800 ms**: accuracy, *r* = 0.18, *p* = 0.21; confidence, *r =* 0.27, *p* = 0.06; **800-1400 ms**: accuracy, *r* = 0.29, *p* < 0.05; confidence, *r* = 0.28, *p* = 0.05). The accuracy and confidence difference scores were themselves correlated, *r* = 0.40, *p* = 0.004.

Next, we considered individual differences in the MDD group alone, again focusing on “Question minus Side” comparisons for words from the mobility task. First we computed Pearson correlations to examine associations between “Question minus Side” accuracy and depressive severity (BDI-II total), anhedonia (MASQ-AD), general distress associated with anxiety (MASQ-GDA), anxious arousal (MASQ-AA), brooding rumination (RRS-Brooding), and sleep disruption (PSQI total). We found negative relationships between accuracy and both general distress due to anxiety (Figure 10A; *r* = -0.42, *p* = 0.04) and anxious arousal (Figure 10B; *r* = -0.48, *p* < 0.02). No other result was significant, but there was a weak relationship with BDI-II total score (*r* = -0.32, *p* = 0.12). To determine whether the relationships with anxiety remained significant after accounting for depressive severity, we computed two hierarchical regressions in which “Question minus Side” accuracy for words from the mobility task served as the criterion variable, BDI-II scores were added in step 1, and either MASQ-GDA or MASQ-AA were entered in step 2. Neither model was significantly improved by addition of the MASQ anxiety measures, and neither MASQ measure reliably predicted accuracy with BDI-II already in the model, although the effect of MASQ-AA was marginally significant (*p* = 0.07). This is not entirely surprising as BDI-II, MASQ-GDA, and MASQ-AA scores were correlated, *r*s > 0.68, *p*s < 0.001, suggesting that they captured overlapping variance in psychological distress. Thus, we were not able to identify an effect of anxiety that was separable from greater depressive severity.

Next we examined relationships with “Question minus Side” ERP difference waves, computed for words from the mobility task, solely in the depressed group. To maximize sensitivity, we did not restrict the analysis to sites that showed group differences (Figure 9). Instead, we extracted data from electrodes that showed condition effects in the MDD group considered alone (Figure 10C). These electrodes were predominantly located over left parietal scalp, and significant clusters were found in all three time windows (400-800 ms, 800-1400 ms, 1400-2000 ms). The mean “Question minus Side” ERP amplitude from these electrodes was correlated across the windows (*r*s > 0.39, *p*s < 0.053), but we did not find strong relationships with accuracy or confidence; the only such finding was a marginal relationship between source accuracy and ERP amplitude from 1400-2000 ms, *r* = 0.39, *p* < 0.06 (all other |*r*|s < 0.26, *p*s > 0.22). However, we found negative correlations between PSQI scores and ERP amplitudes from 400-800 ms (Figure 10D; *r* = -0.48, *p* = 0.02) and 800-1400 ms (Figure 10E; *r* = -0.50, *p* < 0.02).

To confirm that these results did not simply reflect depressive severity, we computed hierarchical regressions with ERP amplitude as the criterion, entering BDI-II and PSQI scores in steps 1 and 2. PSQI predicted ERP amplitude after accounting for BDI-II in both intervals (400-800 ms; β = -0.45, *p* < 0.05; 800-1400 ms; β = -0.49, *p* = 0.03), and adding PSQI improved both models (Δ*R*2s > 0.16, Δ*F*s > 4.5, *p*s < 0.05). Thus, “Question minus Side” ERP amplitude over left parietal scalp in response to words from the mobility task was lowest in depressed adults who reported chronic sleep disruption, and this was statistically distinguisable from greater depressive severity.

[PLEASE INSERT FIGURE 10 ABOUT HERE]

**4. Discussion**

This study yielded three novel findings. First, there was a strong *Cue* x *Task* interaction for source accuracy across participants; conceptual and perceptual retrieval were similarly accurate for words from the mobility task (although this was qualified by the *Group* x *Cue* interaction, discussed below), but conceptual accuracy was substantially worse than perceptual accuracy for words from the animacy task. In other words, conceptual memory was more strongly shaped by the encoding tasks than was perceptual memory. The dependency of conceptual retrieval on encoding was unanticipated, as prior work with this paradigm has collapsed over the encoding tasks (Bergström et al., 2013; Simons et al., 2005a,b), but the results evoke a key finding from behavioral studies of multidimensional source memory: retrieval of any one contextual attribute of an item appears to occur independently of retrieval of any other attribute (Hicks and Starns, 2016; Starns and Hicks, 2005; Vogt and Bröder, 2007).

For instance, across six studies Starns and Hicks manipulated two dimensions of the encoding context, namely font size and word location (Starns and Hicks, 2005) or the gender of faces (on which words were superimposed) and word location (Hicks and Starns, 2016). They then provided cues relevant to one dimension at retrieval—for example, reinstating the font size of some words at test. If the various attributes of the encoded words were directly bound to each another, one would expect these cues to improve not only “within-dimension” retrieval (e.g., memory for font size), but also “across-dimension” retrieval (e.g., memory for word location). However, the six experiments yielded no evidence for across-dimension cueing, with the posterior distribution for this effect squarely centered on zero (see also Vogt and Bröder, 2007). The fact that conceptual retrieval was more sensitive to the encoding tasks than perceptual retrieval is consistent with this prior work, in the sense that encoding influenced retrieval of one attribute (encoding task) while leaving retrieval of another attributes (encoding location) unaffected (for additonal evidence pertinent to encoding manipulations, see Marsh et al., 2004).

We offer two hypotheses to try to explain why conceptual retrieval was more accurate following the mobility vs. animacy task. First, it seems plausible that conceptual encoding was simply more effective during the mobility task. Encoding RT was slower on mobility (vs. animacy) trials, and because both tasks required analysis of the semantic properties of words, we regard longer RTs as evidence of sustained analysis—deeper processing—in the mobility task. This argument is similar to one made by Dobbins and Wagner (2005), who found slower RTs for pleasantness vs. animacy judgments. They suggested that the basis for making pleasantness judgments varies more from item to item than does the basis for making animacy judgments, and thus concluded that the former involved more “sustained conceptual analysis” than the latter. Similarly, the cognitive processes involved in mobility judgments are likely more numerous and varied than those involved in animacy judgments; for example, it seems easier to decide whether or not *“*elm” refers to a living thing than to decide whether or not an “elm” is mobile, as trees sway in the breeze but clearly do not move on their own power. Consequently, Question accuracy may have been better for words from the mobility task simply because participants engaged in deeper processing and thus generated stronger memory traces on those trials relative to animacy trials. The second possibility is closely related: perhaps the similarity of the cognitive processes engaged by both encoding tasks, combined with deeper processing on mobility trials, resulted in heightened interference when words from the animacy task were presented under the Question cue. In other words, when confronted with the more weakly encoded words from the animacy task, participants may have inadvertently retrieved material that seemed consistent with the extended processes engaged by the mobility task, and thus erroneously endorsed that conceptual source.

Whatever the precise psychological mechanism, the data in Figure 8 link Question hits for words from the animacy task to brain activity not typically observed in studies of episodic retrieval. Specifically, the “Question minus Side” contrast for words from the animacy task did not reveal positive ERPs over the left parietal cortex from 400-800 ms (seen in the “Question minus Odd/Even” [Figure 6] and “Question minus Side”/*mobility* [Figure 9] contrasts, as well as many prior studies (Rugg and Curran, 2007)), nor did it reveal an LPN centered over posterior regions (as seen in the “Side minus Odd/Even” [Figure 7] and “Question minus Side”/*mobility* [Figure 9] contrasts, and many prior studies (Johansson and Mecklinger, 2003)). Instead, this contrast revealed broadly distributed, sustained negative-going potentials that appeared to have a fronto-central focus, with a shift towards the left hemisphere in the 400-800 ms and 1400-2000 ms time windows. To our knowledge, this ERP effect has not been previously reported in studies of episodic retrieval and its functional correlates are unclear. However, mediofrontal negativities are often elicited by outcomes that are worse than expected, and it has been suggested that this reflects a signal to increase cognitive control to improve performance (e.g., Potts et al., 2006). The tasks typically used to elicit mediofrontal negativities differ in many respects from the task used here, but because accuracy was relatively poor for animacy words presented under the Question cue, we speculate that the ERPs in Figure 8 may reflect increased cognitive control in an attempt to respond accurately in this difficult condition.

Second, we found surprisingly good performance—few guesses, high accuracy—in the MDD group when words from the mobility task were presented under the Question cue. We view the accuracy boost as consistent with the cognitive initiative framework, because this cell of the design likely offered the most support at encoding and retrieval. As noted earlier, the mobility task appears to have elicited deeper processing than the animacy task at encoding. At retrieval, responses to the Question cue were made more confidently and markedly more slowly than responses to the Side cue. Thus, the results shown in the left panels of Figure 4A and Figure 4B are in line with prior findings: given engaging encoding and retrieval tasks, episodic memory in depressed adults can be surprisingly good. Perhaps the most interesting result is the fact that this behavioral effect was linked to the amplitude of left parietal ERPs (Figure 9) consistently associated with recollection (Rugg and Curran, 2007). Based on this finding, we speculate that whether a given episodic memory task yields better vs. worse performance in depressed vs. healthy adults may hinge on the extent to which this left parietal circuitry—and the core recollection network more broadly (Rugg and Vilberg, 2013)—is engaged. Finally, we note that in depressed adults relative to controls, confidence was significantly lower on Side trials and left parietal ERPs elicited by words from the mobility task were substantially lower under the Side cue. This offers support to the cognitive initiative framework from another quarter, by showing that depressed participants performed worst when cued to retrieve an arbitrarily assigned attribute totally unrelated to the items’ semantic properties—namely, the position of the words on the screen.

Third, in the MDD group the amplitude of the left parietal ERPs was negatively correlated with self-reported sleep disruption over the last month, and this effect remained significant after controlling for depressive severity. We are unaware of prior ERP or imaging data on the relationship between sleep and episodic retrieval in MDD, but this result is conceptually consistent with prior studies of working memory in healthy adults. For instance, an fMRI study examined activation of the fronto-parietal control network as healthy adults performed a working memory task when they were well-rested, and again after 24 and 35 hours of total sleep deprivation (Chee et al., 2006). There was a main effect of deprivation on activation of the superior parietal lobe, with the strongest effects in the left hemisphere: parietal activation was sharply decreased in the deprived vs. the rested state. Moreover, the amplitude of left parietal and left frontal activation at baseline negatively predicted deprivation-induced drops in accuracy, in a manner akin to the “cognitive reserve” hypothesis of aging (Stern, 2012): participants with greater activation in these regions at baseline were least affected by sleep deprivation. Our detection of a negative effect of disrupted sleep on left parietal ERPs in MDD is in line with these prior findings, but we did not find any relationship between sleep and source accuracy or confidence. We expect that such a relationship will be detected in future work, because sleep disruption is extremely common in depression (Tsuno et al., 2005), sleep disruption exerts broad negative effects on range of cognitive functions (Durmer and Dinges, 2005), and there is reason to think that disrupted episodic retrieval in MDD may reflect reduced executive function (Dalgleish et al., 2007). Thus, we speculate that sleep disruption may make a key contribution to impaired episodic retrieval for neutral information in depression. In fact, we suspect that sleep deprivation may prove key to emotional memory biases in MDD as well, since poor sleep appears to impair long-term retention of positive and neutral memories more than negative memories (Sterpenich et al., 2007)—this could contribute to the memory advantage for negative stimuli in adults with MDD. Finally, drowsiness is associated with increased alpha power in the EEG, and alpha is associated with inhibition of cortical function (Jensen and Mazaheri, 2010). Thus, a fascinating topic for future work would be to determine whether sleep disruption in psychiatric groups is linked to cognitive deficits via alpha-induced inhibition of key cortical areas—here, left parietal regions important for source memory.

Although we did not observe group differences when Question and Side hits (collapsed over the encoding tasks) were contrasted to correct responses on Odd/Even trials (Figures 6 and 7), the results of these contrasts were very similar to prior reports. Both contrasts elicited left parietal activity from 400-800 ms, although this was only significant in the Question vs. Odd/Even contrast (the lack of significance in the Side vs. Odd/Even contrast reflects the stringency of the multiple comparisons correction used in our mass univariate analysis, as well as the fact that confidence was consistently lower on Side vs. Question trials—for the link between confidence and recollection during source memory judgments, see Thakral et al., 2015). The parietal effects were followed by very robust LPNs, which were evident over midline posterior scalp in both contrasts, albeit with a more right hemisphere focus for Question vs. Odd/Even, and extended over left frontal scalp in the Question vs. Odd/Even contrast. The midline posterior results are consistent with the aforementioned PMAT framework (Ritchey et al., 2015), which links the retrieval of contextual information in general and spatial information in particular to activation of the retrosplenial cortex and the precuneus. Because the multi-modal study by Bergström and colleagues (2013) linked the LPN to fMRI activation in the precuneus, we interpret the posterior LPN as reflecting process related to contextual retrieval. The nature of those processes is unclear, but candidates include post-retrieval monitoring (Simons et al., 2010), maintenance of retrieved information in an episodic buffer (Baddeley, 2000), direction of attention to the products of the retrieval search (Cabeza et al., 2008), or perhaps the generation or maintenance of visual images elicited by any of these processes (Farah, 1984, 1989). The left frontal LPN seen in the Question vs. Odd/Even contrast may reflect cue elaboration (Han et al., 2012), selection among candidate memories (Bergström et al., 2015), or possibly reinstatement of left frontal activity that supported performance of the encoding tasks in the first place (Thakral et al., 2015). Given the complexity of these processes and the robust evidence of cognitive dysfunction in depression (Snyder, 2013), we expect that depression-related deficits may yet be detectable in future studies. And while we cannot adjudicate between the various candidates in this dataset, we note that because the LPN follows the left parietal effect, it very likely mediates a process (or processes) that act on the products of recollection, and probably does not reflect activity related to initiating retrieval searches (Bergström et al., 2013).

It is important to be aware of limitations associated with the experimental design. First, we cannot conclusively link the ERP results in Figure 9 to accuracy vs. confidence. We are inclined to interpret them as reflecting fluctuation in memory accuracy because the patterns in Figure 9 (bottom panel) and Figure 4 (top left panel) are quite similar, but across the groups the ERPs were weakly correlated with both accuracy and confidence. Furthermore, “Question minus Side” accuracy and confidence difference scores were significantly positively correlated for words from the mobility task, which suggests that cleanly dissociating effects associated with accuracy vs. confidence may not be possible in these data.

Second, we cannot be certain that the *Cue* x *Task* accuracy interaction observed across the groups truly reflects a difference in memory as opposed to response bias. That is, it is possible that participants were simply inclined to respond “mobility task” under the Question whenever they were uncertain, and this—rather than a difference in veridical recall—is what led to the interaction. Unfortunately, we know of no way to distinguish between these possibilities. It is possible to compute *d*’ for responses to the Question and Side cues in this task by arbitrarily calling correct responses to one source (mobility task, right side) “hits” and incorrect endorsements of these sources (i.e., responding “mobility task” to words from the animacy task, or “right” to words presented on the left) “false alarms” (Hicks and Starns, 2016; Slotnick and Dodson, 2005). When we do this, we find a significant *Group* x *Cue* interaction that reflects (non-significantly) higher *d*’ scores for depressed vs. healthy adults under the Question cue and vice versa under the Side cue (analysis not shown). These results are encouraging as they indicate that our data are not confounded with a strong group difference in response bias. Furthermore, the pattern of results is broadly consistent with the accuracy and confidence data presented earlier (i.e., the MDD group shows a Question > Side advantage). However, the need to incorporate both encoding tasks into the *d*’calculation makes it impossible to detect the *Cue* x *Task* interactions that are clearly important here. Thus, separating changes in accuracy from shifts in response bias is a goal for future work on source memory in MDD.

Third, the lack of new items in this paradigm is a problem, although it does confer one advantage: because all the words are “old” at retrieval, there is no reason to suspect that source accuracy is confounded with gross differences in recognition memory, which is frequently the case in designs that require old/new judgments before source decisions are rendered (Murnane and Bayen, 1996). However, the lack of new items contributes to the aforementioned challenges regarding teasing apart accuracy vs. response bias, and it also makes it impossible to determine whether familiarity made any contribution to performance or was affected by MDD. We were aware of this issue at the outset of the study and decided that it was not of critical importance because MDD affects recollection much more strongly than familiarity (Hertel and Milan, 1994; MacQueen et al., 2002), but it would be preferable to include new items in future studies.

A fourth limitation is that, with 24 individuals per group, the power of this study is relatively low. The sample sizes reflect the challenges associated with recruiting unmedicated depressed participants without substantial comorbidity, and we note that they are in line with prior work in this area. However, we are entirely in agreement with the recent emphasis on the need for better-powered studies in psychology and neuroscience (Button et al., 2013), and it is possible that we might have detected stronger negative effects of depression with more participants. Other methodological changes that would likely facilitate the detection of negative effects of depression include the use of less structured encoding and retrieval tasks, as well as the inclusion of emotionally positive stimuli, which would be expected to enhance memory in the controls more than in the depressed adults. Finally, it is important to emphasize again that our *a priori* hypotheses were not supported and thus the ERP results presented in Figures 8 and 9 constitute a post-hoc analysis, which raises the likelihood of Type I errors.

We believe that this line of research is valuable despite these limitations. Much of the early work on memory in MDD was conducted by neuropsychologists who wanted to know how to tell depression from dementia (Burt et al., 1995). But memory deficits in depression deserve to be well-understood for their own sake. The hopelessness that characterizes depression is bound up in negative autobiographical memories, and thus a better understanding of the way in which these memories are encoded, consolidated, and retrieved could yield insight into the pathophysiology of MDD. Furthermore, autobiographical memory retrieval in depression is overly general (Williams et al., 2007), and this has been linked to poor problem-solving in the present and imprecise thinking about the future (Williams et al., 1996). Because cognitive behavioral therapy (CBT) depends on the patient’s ability to recall difficult episodes from their lives and then imagine similar situations unfolding more positively going forward (Holmes et al., 2007), we suspect that there are important but underappreciated links between episodic retrieval, prospection, and the ability to benefit from CBT. Indeed, the new field of “memory therapeutics” reports that enhancing episodic memory—for non-emotional as well as emotional stimuli—leads to meaningful clinical improvements (for review, see Dalgleish and Werner-Seidler, 2014), and it is intriguing that imagining future events depends on parietal circuits (Madore et al., 2016) similar to those shown in Figure 9. Finally, depression is often recurrent as well as debilitating, making it a leading cause of years lived with disability worldwide (Ferrari et al., 2013) that costs the U.S. billions of dollars per year (Greenberg et al., 2015). There is an acute need for better treatments (Fournier et al., 2010), and there is evidence from non-human animals linking the effects of antidepressants to structural changes in brain regions critically implicated in episodic memory (Perera et al., 2007; Santarelli et al., 2003, 2008). Determining whether these findings extend to humans depends on a better understanding of how MDD affects the neural machinery that supports episodic memory. This study takes a small step in this important direction.

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**References**

Airaksinen, E., Larsson, M., Lundberg, I., Forsell, Y. **Cognitive functions in depressive disorders: Evidence from a population-based study**. Psychol. Med. 34 (2004), pp. 83–91.

Baddeley, A. **The episodic buffer: a new component of working memory?** Trends Cogn. Sci. 4 (2000), pp. 417-423.

Beck, A.T., Steer, R.A., Brown, G.K. (1996). **Manual for the Beck depression inventory-II**. The Psychological Corporation, San Antonio, TX.

Bergström, Z.M., Henson, R.N., Taylor, J.R., Simons, J.S.. **Multimodal imaging reveals the spatiotemporal dynamics of recollection**. Neuroimage 68 (2013), pp. 141–153.

Bremner, J.D., Vythilingam, M., Vermetten, E., Vaccarino, V., Charney, D.S. **Deficits in hippocampal and anterior cingulate functioning during verbal declarative memory encoding in midlife major depression**. Am. J. Psychiatry 161 (2004), pp. 637–645.

Bullmore, E.T., Suckling, J., Overmeyer, S., Rabe-Hesketh, S., Taylor, E., Brammer, M.J. **Global, voxel, and cluster tests, by theory and permutation, for a difference between two groups of structural MR images of the brain**. IEEE Trans. Med. Imaging 18 (1999), pp. 32–42.

Burt, D.B., Zembar, M.J., Niederehe, G. **Depression and memory impairment: a meta-analysis of the association, its pattern, and specificity**. Psychol. Bull. 117 (1995), pp. 285–305.

Button, K.S., Ioannidis, J.P., Mokrysz, C., Nosek, B.A., Flint, J., Robinson, E.S., Munafò, M.R. **Power failure: why small sample size undermines the reliability of neuroscience**. Nat. Rev. Neurosci. 14, pp. 365-376.

Buysse, D.J., Reynolds, C.F., Monk, T.H., Berman, S.R., Kupfer, D.J., III, C.F.R., Monk, T.H., Berman, S.R., Kupfer, D.J. **The Pittsburgh Sleep Quality Index: a new instrument for psychiatric practice and research**. Psychiatry Res. 28 (1989), pp. 193–213.

Cabeza, R., Ciaramelli, E., Olson, I.R., Moscovitch, M. **Parietal cortex and episodic memory: an attentional account**. Nat. Rev. Neurosci. 9 (2008), pp. 613-625.

Chee, M.W.L., Chuah, L.Y.M., Venkatraman, V., Chan, W.Y., Philip, P., Dinges, D.F. **Functional imaging of working memory following normal sleep and after 24 and 35 h of sleep deprivation: Correlations of fronto-parietal activation with performance**. Neuroimage 31 (2006), pp. 419–428.

Coltheart, M. **The MRC psycholinguistic database**. Q. J. Exp. Psychol. Sect. A 33 (1981), pp. 497–505.

Cooney, R.E., Joormann, J., Eugène, F., Dennis, E.L., Gotlib, I.H. **Neural correlates of rumination in depression**. Cogn. Affect. Behav. Neurosci. 10 (2010), pp. 470–478.

Cycowicz, Y.M., Friedman, D., Snodgrass, J.G. **Remembering the color of objects: an ERP investigation of source memory**. Cereb. Cortex 11 (2001), pp. 322–334.

Dalgleish, T., Werner-Seidler, A. **Disruptions in autobiographical memory processing in depression and the emergence of memory therapeutics**. Trends Cogn. Sci. 18 (2014), pp. 596–604.

Dalgleish, T., Williams, J.M.G., Golden, A.J., Perkins, N., Barrett, L.F., Barnard, P.J., Yeung, C.A., Murphy, V., Elward, R., Tchanturia, K., Watkins, E. **Reduced specificity of autobiographical memory and depression: the role of executive control**. J. Exp. Psychol. Gen. 136 (2007), pp. 23–42.

Degl’Innocenti, A., Bäckman, L. **Source memory in major depression**. J. Affect. Disord. 54 (1999), pp. 205–209.

Deldin, P.J., Phillips, L.K., Thomas, R.J. **A preliminary study of sleep-disordered breathing in major depressive disorder**. Sleep Med. 7 (2006), pp. 131–139.

Delorme, A., Makeig, S. **EEGLAB:** **An open source toolbox for analysis of single-trial EEG dynamics including independent component analysis**. J. Neurosci. Methods 134 (2004), pp. 9–21.

Dillon, D.G. **The neuroscience of positive memory deficits in depression**. Front. Psychol. 6 (2015), 1295.

Dillon, D.G., Dobbins, I.G., Pizzagalli, D.A. **Weak reward source memory in depression reflects blunted activation of VTA/SN and parahippocampus**. Soc. Cogn. Affect. Neurosci. 9 (2014), pp. 1576–1583.

Dillon, D.G., Pizzagalli, D.A. **Evidence of successful modulation of brain activation and subjective experience during reappraisal of negative emotion in unmedicated depression**. Psychiatry Res. - Neuroimaging 212 (2013), pp. 99-107.

Dobbins, I.G., Wagner, A.D. **Domain-general and domain-sensitive prefrontal mechanisms for recollecting events and detecting novelty**. Cereb. Cortex 15 (2005), pp. 1768–1778.

Durmer, J.S., Dinges, D.F. **Neurocognitive consequences of sleep deprivation**. Semin. Neurol. 25 (2005), pp. 117–129.

Eichenbaum, H., Yonelinas, A.P., Ranganath, C. **The medial temporal lobe and recognition memory**. Annu. Rev. Neurosci. 30 (2007), pp. 123–152.

Farah, M.J. **The neurological basis of mental imagery: a componential analysis**. Cognition 18 (1984), pp. 245-272.

Farah, M.J. **The neural basis of mental imagery**. Trends Neurosci. 12 (1989), pp. 395-399.

Ferrari, A.J., Charlson, F.J., Norman, R.E., Patten, S.B., Freedman, G., Murray, C.J.L., Vos, T., Whiteford, H.A. **Burden of depressive disorders by country, sex, age, and year: findings from the Global Burden of Disease Study 2010**. PLoS Med. 10 (2013). e1001547

Fournier, J.C., DeRubeis, R.J., Hollon, S.D., Dimidjian, S., Amsterdam, J.D., Shelton, R.C., Fawcett, J. **Antidepressant drug effects and depression severity: a patient-level meta-analysis**. JAMA 303 (2010), pp. 47–53.

Friston, K.J., Holmes, A.P., Worsley, K.J., Poline, J.-P., Frith, C.D., Frackowiak, R.S.J. **Statistical parametric maps in functional imaging: A general linear approach**. Hum. Brain Mapp. 2 (1995), pp. 189–210.

Greenberg, P.E., Fournier, A.-A., Sisitsky, T., Pike, C.T., Kessler, R.C. **The economic burden of adults with major depressive disorder in the United States (2005 and 2010)**. J. Clin. Psychiatry 76 (2015), pp. 155–162.

Groppe, D.M., Urbach, T.P., Kutas, M. **Mass univariate analysis of event-related brain potentials/fields I: A critical tutorial review**. Psychophysiology 48 (2011a), pp. 1711-1725

Groppe, D.M., Urbach, T.P., Kutas, M. **Mass univariate analysis of event-related brain potentials/fields II: Simulation studies**. Psychophysiology 48 (2011b), pp. 1726–1737.

Hamilton, J.P., Gotlib, I.H. **Neural substrates of increased memory sensitivity for negative stimuli in major depression**. Biol. Psychiatry 63 (2008), pp. 1155–1162.

Han, S., OʼConnor, A.R., Eslick, A.N., Dobbins, I.G. **The role of left ventrolateral prefrontal cortex during episodic decisions: semantic elaboration or resolution of episodic interference?** J. Cogn. Neurosci. 24 (2012), pp. 223-234.

Hertel, P.T. **On the contributions of deficient cognitive control to memory impairments in depression**. Cogn. Emot. 11 (1997), pp. 569–583.

Hertel, P.T., Hardin, T.S. **Remembering with and without awareness in a depressed mood: Evidence of deficits in initiative**. J. Exp. Psychol. Gen. 119 (1990), pp. 45–59.

Hertel, P.T., Milan, S. **Depressive deficits in recognition: Dissociation of recollection and familiarity**. J. Abnorm. Psychol. 103 (1994), pp. 736–742.

Hertel, P.T., Rude, S.S. **Depressive deficits in memory: focusing attention improves subsequent recall**. J. Exp. Psychol. Gen. 120 (1991), pp. 301–309.

Hertel, P.T. **Relation between rumination and impaired memory in dysphoric moods**. J. Abnorm. Psychol. 107 (1998), pp. 166–172.

Hicks, J.L., Starns, J.J. **Successful cuing of gender source memory does not improve location source memory**. Mem. Cognit. 44 (2016), 650–659.

Holdnack, H.A. (2001). **Wechsler Test of Adult Reading: WTAR**. The Psychological Corporation, San Antonio, TX.

Holmes, E.A., Arntz, A., Smucker, M.R. **Imagery rescripting in cognitive behaviour therapy: Images, treatment techniques and outcomes**. J. Behav. Ther. Exp. Psychiatry 38 (2007), pp. 297–305.

Jensen, O., Mazaheri, A. **Shaping functional architecture by oscillatory alpha activity: gating by inhibition**. Front. Hum. Neurosci. 4 (2010), 186.

Johansson, M., Mecklinger, A. **The late posterior negativity in ERP studies of episodic memory: action monitoring and retrieval of attribute conjunctions**. Biol. Psychol. 64 (2003), pp. 91–117.

Johnson, M.K., Hashtroudi, S., Lindsay, D.S. **Source monitoring**. Psychol. Bull. 114 (1993), pp. 3-28.

Lopez-Calderon, J., Luck, S.J. **ERPLAB: an open-source toolbox for the analysis of event-related potentials**. Front. Hum. Neurosci. 8 (2014), 213.

Luck, S.J. (2014). **An introduction to the event-related potential technique, 2nd ed**. MIT Press, Cambridge, MA.

MacQueen, G., Frodl, T. **The hippocampus in major depression: evidence for the convergence of the bench and bedside in psychiatric research?** Mol. Psychiatry 16 (2014), pp. 252–264.

MacQueen, G.M., Galway, T.M., Hay, J., Young, L.T., Joffe, R.T. **Recollection memory deficits in patients with major depressive disorder predicted by past depressions but not current mood state or treatment status**. Psychol. Med. 32 (2002), pp. 251–258.

Madore, K.P., Szpunar, K.K., Addis, D.R., Schacter, D.L. **Episodic specificity induction impacts activity in a core brain network during construction of imagined future experiences**. Proc. Natl. Acad. Sci. U. S. A. 113 (2016), pp. 10696–10701.

Marsh, R.L., Hicks, J.L., Cook, G.I. **Focused attention on one contextual attribute does not reduce source memory for a different attribute**. Memory 12 (2004), pp. 183–192.

Matt, G.E., Vazquez, C., Campbell, W.K. **Mood-congruent recall of affectively toned stimuli: A meta-analytic review**. Clin. Psychol. Rev. 12 (1992), pp. 227–255.

McEwen, B.S. **Sleep deprivation as a neurobiologic and physiologic stressor: allostasis and allostatic load**. Metabolism. 55 (2006), pp. S20-S23.

Mecklinger, A., Johansson, M., Parra, M., Hanslmayr, S. **Source-retrieval requirements influence late ERP and EEG memory effects**. Brain Res. 1172 (2007), pp. 110–123.

Mollison, M. V, Curran, T. **Familiarity in source memory**. Neuropsychologia 50 (2012), pp. 2546–2565.

Murnane, K., Bayen, U.J. **An evaluation of empirical measures of source identification**. Mem. Cognit. 24 (1996), pp. 417–428.

Nolen-Hoeksema, S. **Responses to depression and their effects on the duration of depressive episodes**. J. Abnorm. Psychol. 100 (1991), pp. 569–582.

Peirce, J.W. **Generating stimuli for neuroscience using PsychoPy**. Front. Neuroinform. 2 (2009), 10.

Perera, T.D., Coplan, J.D., Lisanby, S.H., Lipira, C.M., Arif, M., Carpio, C., Spitzer, G., Santarelli, L., Scharf, B., Hen, R., Rosoklija, G., Sackeim, H. A, Dwork, A.J. **Antidepressant-induced neurogenesis in the hippocampus of adult nonhuman primates**. J. Neurosci. 27 (2007), pp. 4894–4901.

Potts, G.F., Martin, L.E., Burton, P., Montague, P.R. **When things are better or worse than expected: the medial frontal cortex and the allocation of processing resources**. J. Cogn. Neurosci. 18 (2006), pp. 1112–1119.

R Core Team (2015). **R: A Language and Environment for Statistical Computing. R Foundation for Statistical Computing**, Vienna, Austria.

Reitman, J.S., Higman, B., Lifson, A., Rosenblum, J. **Without surreptitious rehearsal, information in short-term memory decays I**. J. Verbal Learning Verbal Behav. 13 (1974), pp. 365–377.

Ritchey, M., Libby, L.A., Ranganath, C. **Cortico-hippocampal systems involved in memory and cognition: The PMAT framework**. Prog. Brain Res. 219 (2015), pp. 45–64.

Rock, P.L., Roiser, J.P., Riedel, W.J., Blackwell, A.D. **Cognitive impairment in depression: a systematic review and meta-analysis**. Psychol. Med. 44 (2014), pp. 2029–2040.

Rugg, M.D., Curran, T. **Event-related potentials and recognition memory**. Trends Cogn. Sci. 11 (2007), pp. 251–257.

Rugg, M.D., Vilberg, K.L. **Brain networks underlying episodic memory retrieval**. Curr. Opin. Neurobiol. 23 (2013), pp. 255–260.

Santarelli, L., Saxe, M., Gross, C., Surget, A., Dulawa, S., Weisstaub, N., Lee, J., Duman, R., Arancio, O., Santarelli, L., Saxe, M., Gross, C., Surget, A., Battaglia, F., Dulawa, S., Weisstaub, N., Lee, J., Duman, R., Arancio, O., Beizung, C., Hen, R., Belzung, C., Hen, R. **Requirement of hippocampal neurogenesis for the behavioral effects of antidepressants**. Science 301 (2003), pp. 805–809.

Sheehan, D. V., Lecrubier, Y., Sheehan, K.H., Amorim, P., Janavs, J., Weiller, E., Hergueta, T., Baker, R., Dunbar, G.C. **The Mini-International Neuropsychiatric Interview (M.I.N.I.): The development and validation of a structured diagnostic psychiatric interview for DSM-IV and ICD-10**. J. Clin. Psychiatry 59 Suppl 20 (1998), pp. 22–33.

Simons, J.S., Gilbert, S.J., Owen, A.M., Fletcher, P.C., Burgess, P.W. **Distinct roles for lateral and medial anterior prefrontal cortex in contextual recollection**. J. Neurophysiol 94 (2005a), pp. 813–820.

Simons, J.S., Owen, A.M., Fletcher, P.C., Burgess, P.W. **Anterior prefrontal cortex and the recollection of contextual information**. Neuropsychologia 43 (2005b), pp. 1774–1783.

Simons, J.S., Peers, P.V., Mazuz, Y.S., Berryhill, M.E., Olson, I.R. **Dissociation between memory accuracy and memory confidence following bilateral parietal lesions**. Cereb. Cortex (2010), pp. 479-485.

Singmann, H., Bolker, B., Westfall, J., Aust, F. **afex: Analysis of Factorial Experiments**. R package version 0.16-1. (2016), https://CRAN.R-project.org/package=afex.

Slotnick, S.D., Dodson, C.S. **Support for a continuous (single-process) model of recognition memory and source memory**. Mem. Cognit. 33 (2005), pp. 151–170.

Snyder, H.R. **Major depressive disorder is associated with broad impairments on neuropsychological measures of executive function: a meta-analysis and review**. Psychol. Bull. 139 (2013), pp. 81-132.

Song, J., Davey, C., Poulsen, C., Luu, P., Turovets, S., Anderson, E., Li, K., Tucker, D. **EEG source localization: Sensor density and head surface coverage**. J. Neurosci. Methods 256 (2015), pp. 9–21.

Starns, J.J., Hicks, J.L. **Source dimensions are retrieved independently in multidimensional monitoring tasks**. J. Exp. Psychol. Learn. Mem. Cogn. 31 (2005), pp. 1213–1220.

Steffens, D.C., Otey, E., Alexopoulos, G.S., Butters, M.A., Cuthbert, B., Ganguli, M., Geda, Y.E., Hendrie, H.C., Krishnan, R.R., Kumar, A., Lopez, O.L., Lyketsos, C.G., Mast, B.T., Morris, J.C., Norton, M.C., Peavy, G.M., Petersen, R.C., Reynolds, C.F., Salloway, S., Welsh-Bohmer, K.A., Yesavage, J. **Perspectives on depression, mild cognitive impairment, and cognitive decline**. Arch. Gen. Psychiatry 63 (2006), pp. 130-138.

Stern, Y. **Cognitive reserve in ageing and Alzheimer’s disease**. Lancet Neurol. 11 (2012), pp. 1006-1012.

Sterpenich, V., Albouy, G., Boly, M., Vandewalle, G., Darsaud, A., Balteau, E., Dang-Vu, T.T., Desseilles, M., D’Argembeau, A., Gais, S., Rauchs, G., Schabus, M., Degueldre, C., Luxen, A., Collette, F., Maquet, P. **Sleep-related hippocampo-cortical interplay during emotional memory recollection**. PLoS Biol. 5 (2007), e282.

Thakral, P.P., Wang, T.H., Rugg, M.D. **Cortical reinstatement and the confidence and accuracy of source memory**. Neuroimage 109 (2015), pp. 118-129.

Treadway, M.T., Waskom, M.L., Dillon, D.G., Holmes, A.J., Park, M.T.M., Chakravarty, M.M., Dutra, S.J., Polli, F.E., Iosifescu, D.V., Fava, M., Gabrieli, J.D.E., Pizzagalli, D.A. **Illness progression, recent stress, and morphometry of hippocampal subfields and medial prefrontal cortex in major depression**. Biol. Psychiatry 77 (2015), pp. 285-294.

Treynor, W., Gonzalez, R., Nolen-Hoeksema, S. **Rumination reconsidered: a psychometric analysis**. Cognit. Ther. Res. 27 (2003), pp. 247–259.

Tsuno, N., Besset, A., Ritchie, K. **Sleep and depression**. J. Clin. Psychiatry 66 (2005), pp. 1254-1269.

Vogt, V., Bröder, A.. **Independent retrieval of source dimensions: an extension of results by Starns and Hicks (2005) and a comment on the ACSIM measure**. J. Exp. Psychol. Learn. Mem. Cogn. 33 (2007), pp. 443–450.

Watson, D., Weber, K., Assenheimer, J.S., Clark, L.A., Strauss, M.E., McCormick, R.A. **Testing a tripartite model: I. Evaluating the convergent and discriminant validity of anxiety and depression symptom scales**. J. Abnorm. Psychol. 104 (1995), pp. 3–14.

Williams, J.M., Ellis, N.C., Tyers, C., Healy, H., Rose, G., MacLeod, A.K. **The specificity of autobiographical memory and imageability of the future**. Mem. Cognit. 24 (1996), pp. 116–125.

Williams, J.M.G., Barnhofer, T., Crane, C., Herman, D., Raes, F., Watkins, E., Dalgleish, T. **Autobiographical memory specificity and emotional disorder**. Psychol. Bull. 133 (2007), pp. 122–148.

Wulff, K., Gatti, S., Wettstein, J.G., Foster, R.G. **Sleep and circadian rhythm disruption in psychiatric and neurodegenerative disease**. Nat. Rev. Neurosci. 11 (2010), pp. 589–599.

Zakzanis, K.K., Leach, L., Kaplan, E. **On the nature and pattern of neurocognitive function in major depressive disorder**. Neuropsychiatry. Neuropsychol. Behav. Neurol. 11 (1998), pp. 111–119.

**Figure Captions**

*Figure 1*. Encoding (left) and recognition (right) trial structures. Encoding trials began with three centrally presented arrows pointing to the side on which the word would appear. The encoding task was presented next, either “living or non-living?” (animacy judgment) or “mobile or immobile?” (mobility judgment, not shown). Finally, a word was presented directly above the encoding question; participants had 3500 ms to respond. Retrieval trials began with presentation of one of three cues (“Side”, “Question”, or “Odd/Even”). After a 1000 ms delay, a word was presented. On Side and Question trials, the word came from the immediately preceding encoding block, while on Odd/Even trials the word was a numeral (e.g., “seventy-seven”). Finally, a response screen was presented and persisted until the participant responded or 10 seconds had elapsed. The response options for a Side trial are displayed. On Question trials, “left” and “right” were replaced with “living/non-living” and “mobile/immobile”, respectively; on Odd/Even trials they were replaced with “odd” and “even”.

*Figure 2*. Encoding (A) response accuracy and (B) correct response time (RT). Responses were slower and less accurate in the mobility task vs. the animacy task. There were no group differences, but data from controls and depressed participants are shown separately for comparison. Error bars = SEM.

*Figure 3*. Mean number of guesses by group, encoding task, and retrieval cue. Data are from 18 depressed and 13 healthy participants who guessed at least once in every condition. There was a main effect of *Task,* as all participants guessed less for words from the mobility (left panel) vs. the animacy task (right panel), and a main effect of *Cue*, as participants guessed less for words shown under the Question cue (blue bars) vs. the Side cue (green bars). In the MDD group, the cue effect on guessing was pronounced for words from the mobility task. Error bars = SEM, \**p* = 0.004.

*Figure 4*. Source memory accuracy (A) under Question (blue bars) and Side (green bars) cues, and for (B) Question minus Side difference scores. In both panels, the left column shows data for words from the mobility task, and the right panel shows data for words from the animacy task. There was a *Group* x *Cue* interaction for words from the mobility task, because the MDD group showed better memory under the Question vs. Side cue but the controls did not. By contrast, the animacy task was characterized by a main effect of *Cue*, as both groups show lower accuracy under Question vs. Side. Bar heights correspond to mean, error bars = SEM, \* *p* = 0.019.

*Figure 5.* (A) The percentage of high confidence responses as a function of group, cue, and task. All participants were more confident when responding to words from the mobility task (left panel) vs. the animacy task (right panel) and when responding to the Question cue (blue bars) vs. the Side cue (green bars). Depressed adults were less confident than controls in response to the Side cue but not the Question cue. Exploratory *t*-tests revealed thatdepressed adults were significantly more confident when responding to the Question vs. Side cue for words from both encoding tasks. (B) Mean correct RT data; all participants responded more slowly to the Question cue (blue bars) vs. the Side cue (green bars). Error bars = SEM, \**p*s < 0.001.

*Figure 6*. Topographies show mass univariate analysis of *Question* minus *Odd/Even* difference waves, from 400-800 ms (top), 800-1400 ms (middle), and 1400-2000 ms (bottom). The data are collapsed across groups as there were no significant between-group differences. On the topographies, electrodes in significant clusters are marked with white circles. The electrode that showed the strongest condition effect in each cluster is marked in red, and waveforms from that electrode are plotted separately for each condition, with the time window shaded in gray. Electrode numbers (e.g., “e109”) give the position on the EGI cap—see the Supplement for a complete map.

*Figure 7*. Topographies show mass univariate analysis of *Side* minus *Odd/Even* difference waves, from 400-800 ms (top), 800-1400 ms (middle), and 1400-2000 ms (bottom). The data are collapsed across groups as there were no significant between-group differences. On the topographies, electrodes in significant clusters are marked with white circles. The electrode that showed the strongest condition effect in each cluster is marked in red, and waveforms from that electrode are plotted separately for each condition, with the time window shaded in gray. Electrode numbers (e.g., “e83”) give the position on the EGI cap—see the Supplement for a complete map. No significant differences between the conditions were found from 400-800 ms.

*Figure 8*. Topographies show mass univariate analysis of *Question* minus *Side* difference waves, for words from the animacy task, from 400-800 ms (top), 800-1400 ms (middle), and 1400-2000 ms (bottom). The data are collapsed across groups as there were no significant between-group differences. On the topographies, electrodes in significant clusters are marked with white circles. The electrode that showed the strongest condition effect in each cluster is marked in red, and waveforms from that electrode are plotted separately for each condition, with the time window shaded in gray. Electrode numbers (e.g., “e16”) give the position on the EGI cap—see the Supplement for a complete map. An orange line separates the two clusters identified from 800-1400 ms.

*Figure 9*. Topographies show group differences in the mass univariate analysis of *Question* minus *Side* difference waves, for words from the mobility task, from 400-800 ms (top) and 800-1400 ms (middle); bottom panel shows the data averaged over all significant electrodes in each cluster, in each time window. On the topographies, electrodes in significant clusters are marked with white circles. The electrode that showed the strongest condition effect in each cluster window is marked in red, and waveforms from that electrode are plotted separately for each condition, with the relevant time window shaded in gray. Electrode numbers (e.g., “e16”) give the position on the EGI cap. The data in each time window were characterized by *Group* x *Cue* interactions: depressed and healthy adults generated similar responses to the Question cue, but depressed adults generated significantly weaker responses to the Side cue (\**p* < 0.025).

*Figure 10*. In the MDD group, there was a negative correlation between “Question minus Side” accuracy difference scores for words from the mobility task and (A) general distress associated with anxiety (MASQ-GDA) and (B) anxious arousal (MASQ-AA). (C) Topographic maps of Question minus Side differences for words from the mobility task in the MDD group considered alone, with electrodes that showed significant condition effects highlighted in white. There were negative relationships between PSQI scores and Question minus Side ERP amplitudes from these electrodes in the (D) 400-800 ms and (E) 800-1400 ms time windows.