

The Eyes Have It: Hippocampal Activity Predicts Expression of Memory in Eye Movements

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DOI 10.1016/j.neuron.2009.08.025

SUMMARY

Although there is widespread agreement that the hippocampus is critical for explicit episodic memory retrieval, it is controversial whether this region can also support indirect expressions of relational memory when explicit retrieval fails. Here, using functional magnetic resonance imaging (fMRI) with concurrent indirect, eye-movement-based memory measures, we obtained evidence that hippocampal activity predicted expressions of relational memory in subsequent patterns of viewing, even when explicit, conscious retrieval failed. Additionally, activity in the lateral prefrontal cortex and functional connectivity between the hippocampus and prefrontal cortex were greater for correct than for incorrect trials. Together, these results suggest that hippocampal activity can support the expression of relational memory even when explicit retrieval fails and that recruitment of a broader cortical network may be required to support explicit associative recognition.

INTRODUCTION

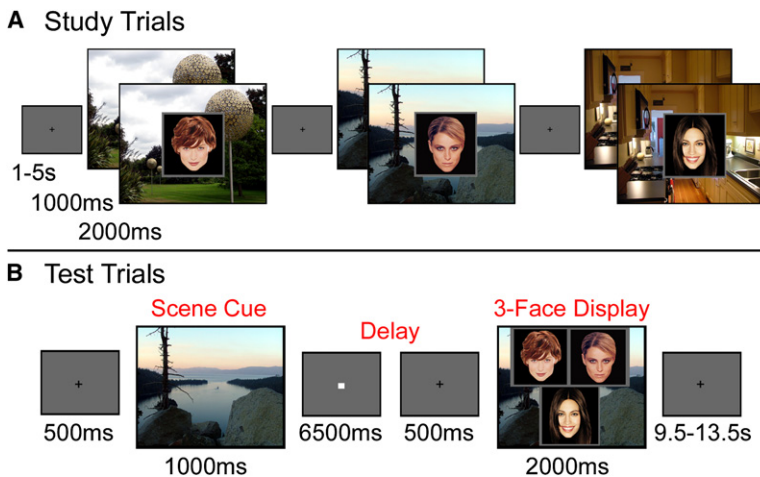
Considerable evidence indicates that the hippocampus and adjacent medial temporal lobe (MTL) cortical structures support long-term declarative memory (Cohen and Squire, 1980; Squire et al., 2004). Several theories implicate these structures specifically in *conscious* retrieval of past events and experiences (e.g., Moscovitch, 1995; Tulving and Schacter, 1990), with particular import placed on the role of the hippocampus in conscious recollection (Aggleton and Brown, 1999; Yonelinas, 2002). An alternative view points to a critical role for the hippocampus in the encoding and retrieval of memories for arbitrary relationships among items that co-occur in the context of some scene or event (Eichenbaum et al., 1994). In general, the relational memory theory is compatible with other accounts of MTL function, as conscious recollection likely depends on the ability to encode, and subsequently retrieve, arbitrary inter-item or item-context relationships (Davachi, 2006; Eichenbaum et al., 2007). However, one area where these theories diverge concerns the role of the hippocampus in the expression of relational memory, even in the absence of awareness. Whereas some theories propose that relationally bound memory representations, supported by the hippocampus, can be expressed even when explicit reports

fail (Eichenbaum et al., 1994; Eichenbaum, 1999), others emphasize the tight link between hippocampal function and conscious retrieval of past events (Aggleton and Brown, 1999; Moscovitch, 1995; Squire et al., 2004; Tulving and Schacter, 1990; Yonelinas, 2002).

Findings from recent experiments suggest that relational memory may be evident in patterns of eye movements even when conscious recollection fails. In these experiments, participants study realistic scenes and are subsequently tested with scenes that are repeated exactly as they were studied and scenes that have been systematically manipulated. Participants typically fixate disproportionately on regions of scenes that have been manipulated, suggesting that memory for the original item-location relationships has modulated viewing patterns (e.g., Hayhoe et al., 1998; Henderson and Hollingworth, 2003; Ryan et al., 2000; Smith et al., 2006). Critically, these eye movement-based relational memory effects have been documented even when participants fail to explicitly detect scene changes (e.g., Hayhoe et al., 1998; Henderson and Hollingworth, 2003; Ryan et al., 2000), suggesting that eye-movement measures can be used to address questions about hippocampal involvement in relational memory retrieval even when overt behavioral reports are incorrect.

Other paradigms have been used to demonstrate that memory can rapidly influence eye-movement behavior (Hannula et al., 2007; Holm et al., 2008; Ryan et al., 2007) and that eye-movement-based memory effects can occur far in advance of explicit recognition (Hannula et al., 2007; Holm et al., 2008). For example, in one study (Hannula et al., 2007), eye movements were monitored during an associative recognition test in which a previously studied scene was presented (“scene cue”), and then three previously studied faces were superimposed on that scene (“test display”). It was hypothesized that the scene cue would elicit expectancies about the face with which it was paired during the study trials, and consistent with this prediction, eye movements were drawn disproportionately to the associated face just 500–750 ms after presentation of the test display. The rapid onset of this effect is notable considering that the position of the associated face could not be predicted and that 500–750 ms is only enough time to permit at most two or three fixations. Furthermore, disproportionate viewing occurred over a second in advance of overt recognition, which suggests that the effect of relational memory on eye-movement behavior might have preceded conscious identification of the match.

The results described above suggest that eye movements can be used to index relational memory retrieval prior to, and possibly even in the absence of, awareness. Accordingly, in the present

**Figure 1. Experimental Paradigm**

(A) Illustration of study trial events. (B) Illustration of a single-test trial.

experiment, we used fMRI with concurrent eye tracking to test whether activity in the hippocampus and/or other MTL regions would be correlated with eye-movement-based relational memory measures even when explicit recognition has failed. Participants in this experiment studied several face-scene pairs, and on each test trial, they were presented with a studied scene, followed by a brief delay, and finally presentation of three studied faces superimposed on that scene (see Figure 1). Critically, one of the faces had been paired with the scene during the study phase (henceforth referred to as the “matching face”), whereas the other two had been paired with different scenes. We expected that presentation of the scene cue would prompt retrieval of the associated face, resulting in increased viewing of that face when the test display was presented (Hannula et al., 2007). The proportion of time spent viewing the matching face was used as an indirect, eye-movement-based measure of relational memory retrieval. We expected that activity in the hippocampus following the scene cue would predict subsequent expression of relational memory in eye-movement behavior, even when conscious recollection failed.

RESULTS

Behavioral Performance: Associative Recognition Accuracy

Participants made accurate responses on 62.29% (SD = 11.10%) of the trials, made incorrect responses on 25.3% (SD = 12.55%) of the trials, and responded “don’t know” on 12.4% (SD = 10.23%) of the trials. Response times were faster for correct (2110.17 ms; SD = 630.80) than for incorrect (2671.73 ms; SD = 850.71) trials, $t(13) = 4.35$, $p < .001$.

Memory for Face-Scene Relationships Is Evident in Eye-Movement Behavior

It was predicted that the scene cue would elicit relational memory retrieval and that this would manifest as rapid, disproportionate viewing of the matching face. Such an effect could not be supported by simple influences of item familiarity, because all three faces in each test display had been seen during the study trials. However, it is reasonable to suppose that participants might

spend more time fixating any face that happened to be selected, even those selected in error. To account for this possibility, we examined whether participants spent more time viewing correctly identified matching faces than faces selected incorrectly. A repeated-measures ANOVA that examined viewing-time data as a function of face type (match, selected) and time bin (0–500, 500–1000, 1000–1500, and 1500–2000 ms) revealed that more time was spent viewing correctly identified matching faces ($M = 0.48$; $SD = 0.08$) than selected faces ($M = 0.40$; $SD = 0.04$), $F(1,13) = 10.88$, $p < 0.01$.

Consistent with previous results (Hannula et al., 2007), disproportionate viewing of matching faces emerged 500–1000ms after the three-face test display was presented ($t(13) = 3.90$, Bonferroni corrected $p < .01$; see Figure 2). These results confirm the rapid influence of relational memory on eye movement behavior, over and above any simple effect of response intention or execution.

MTL Activity during the Scene Cue Predicts Disproportionate Viewing of Matching Faces

Initial fMRI analyses examined the relationship between MTL activity and eye-movement behavior by contrasting trials according to whether participants spent a disproportionate amount of time viewing the matching face (“DPM” trials) or a disproportionate amount of time viewing one of the nonmatching faces (“DPNM” trials). The criterion for disproportionate viewing in this analysis was that the proportion of time spent viewing one face had to exceed the proportion of time spent viewing the remaining two faces by at least 10% (see Supplemental Data for details). We reasoned that, on DPM trials, participants had successfully retrieved information about the previously studied face-scene relationship that was sufficient to influence subsequent eye-movement behavior, whereas this did not occur on DPNM trials (Figure S2 illustrates the time course of these viewing effects); importantly, response times to DPM (2296.66 ms, $SD = 693.96$) and DPNM (2583.81 ms, $SD = 825.43$) trials were not significantly different, $t(13) = 1.50$, $p = 0.16$.

Based on the idea that the hippocampus and adjacent MTL cortical structures are critical for relational memory retrieval, we predicted that activity in these regions during the scene cue would be greater for DPM than for DPNM trials. Consistent with this prediction, BOLD signal was greater for DPM than for DPNM trials in two regions of the right hippocampus (anterior local maximum at $x = 30$, $y = -12$, $z = -24$; $t(13) = 4.06$; posterior local maxima at $x = 24$, $y = -27$, $z = -9$; $t(13) = 3.94$), the right parahippocampal cortex (local maxima at $x = 30$, $y = -27$, $z = -18$; $t(13) = 3.46$), and bilaterally in anterior regions of the parahippocampal gyrus, which likely correspond to the perirhinal cortex (Insausti et al., 1998; left local maxima at $x = -33$, $y = -9$, $z = -36$; $t(13) = 4.21$; right local maxima at $x = 33$, $y = -18$,

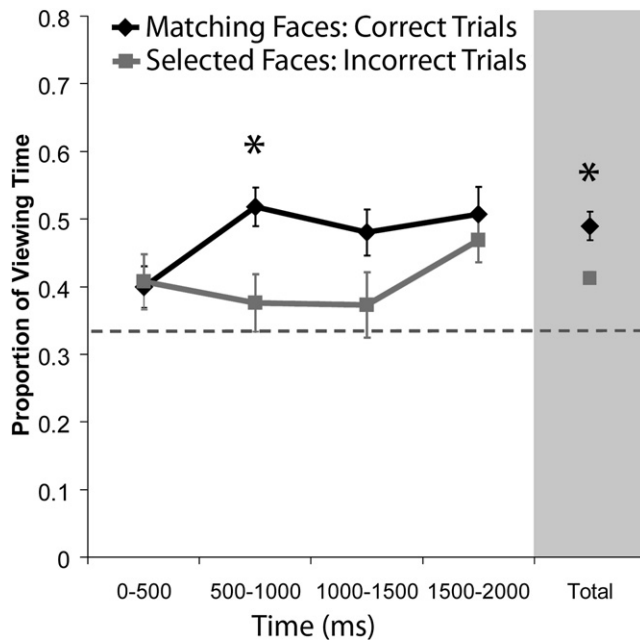


Figure 2. Relational Memory Rapidly Influences Eye-Movement Behavior

Mean proportion of viewing time allocated to the matching face (correct trials) and to selected faces (incorrect trials). Viewing time measures are plotted in successive 500 ms time bins, starting with the onset of the three-face test display. More time was spent viewing correctly identified matching faces than faces that were selected on incorrect trials just 500–1000 ms after the three-face display was presented. The proportion of total viewing time allocated to each face collapsed across the entire 2 s test trial is also illustrated. Standard error bars are plotted around the means; the dashed line represents chance viewing.

$z = -30$; $t(13) = 5.31$). Representative trial-averaged time courses are presented in Figure 3A.

Because response accuracy was greater for DPM trials ($M = 83.30\%$, $SD = 3.56$) than for DPNM trials ($M = 35.20\%$, $SD = 3.80$), it could be argued that correlations between MTL activity and eye movements simply reflected explicit relational memory retrieval. Accordingly, we performed follow-up fMRI analyses to more specifically test whether MTL activity might index eye-movement-based relational memory effects even on trials for which overt recognition failed. In these analyses, we focused specifically on trials for which participants failed to identify the matching face. A median split, based on the proportion of total viewing time directed to the matching face, was used to separately bin trials that were associated with relatively high or low viewing of that face (Figure S3 illustrates the time course of these viewing effects). A mapwise analysis in which activity during the scene cue was contrasted between incorrect high- and incorrect low-viewing trials revealed suprathreshold voxels in bilateral regions of the hippocampus (left local maxima: $x = -24$, $y = -30$, $z = 6$; $t(13) = 5.39$; right local maxima: $x = 27$, $y = -27$, $z = -6$; $t(13) = 4.14$; see Figure 3B). This result implicates the hippocampus in retrieval of information about previously studied face-scene relationships that is sufficient to

influence eye-movement behavior even when explicit recognition has failed.

Perirhinal and Prefrontal Activity during the Scene Cue Predicts Accuracy

The next fMRI analysis examined MTL activity during the scene cue as a function of accuracy, irrespective of eye-movement behavior. Activity during the scene cue was greater for correct than for incorrect trials in a region of left perirhinal cortex, with an activation peak close to the one observed for the disproportionate viewing contrast (local maxima at $x = -21$, $y = 0$, $z = -36$; $t(13) = 5.65$). Surprisingly, there were no suprathreshold activity differences in the hippocampus or the parahippocampal cortex during any part of the test trial. Outside of the MTL, however, several cortical regions (see Table S1) showed increased activity during correct, as compared with incorrect trials, including left dorsolateral prefrontal cortex (DLPFC: local maxima at $x = -48$, $y = 27$, $z = 30$; $t(13) = 5.36$) and left ventrolateral prefrontal cortex (VLPFC: local maxima at $x = -48$, $y = 42$, $z = 0$; $t(13) = 8.38$; see Figure 4A). Results from several studies suggest that these prefrontal regions may implement control processes that support explicit memory attributions (e.g., Dobbins and Sanghooon, 2006; Ranganath et al., 2000; for a review see Fletcher and Henson, 2001).

In order to determine whether PFC activity was also correlated with relational memory as expressed indirectly in eye-movement behavior, parameter estimates for DPM and DPNM trials were extracted from each prefrontal ROI. Following presentation of the scene cue, activity in both regions was greater for DPM than for DPNM trials (left DLPFC: $t(13) = 2.70$, $p < 0.05$; left VLPFC: $t(13) = 2.34$, $p < 0.05$); local maxima identified in the direct contrast of DPM versus DPNM trials are summarized in Table S2. As indicated earlier, however, eye movements were strongly associated with behavioral response accuracy, so this result does not necessarily indicate whether activity in these ROIs was predictive of eye-movement behavior even when recognition failed. To test this possibility, parameter estimates were extracted from the prefrontal ROIs for incorrect trials on which viewing of the match was high versus low. Unlike what was observed in the hippocampus, activity in these ROIs did not differentiate between incorrect high- and low-viewing trials (all t values ≤ 1.87 , all p values > 0.05); local maxima identified in the direct contrast of incorrect high- versus incorrect low-viewing trials are summarized in Table S3.

Functional Connectivity between Hippocampus and PFC Is Increased during Accurate Associative Recognition

Results described above are consistent with the possibility that the hippocampus supports recovery of relational memory and that this information may be communicated to prefrontal regions in order to guide overt decision behavior. If this view is correct, then one might expect increased functional connectivity between the prefrontal regions and the MTL for correct, as compared to incorrect trials. To test this prediction, we ran functional connectivity analyses using the prefrontal ROIs identified in the accuracy contrast as seed regions. Estimates of activity during each phase of each trial were separately averaged within the seed regions for correct and incorrect trials, and these

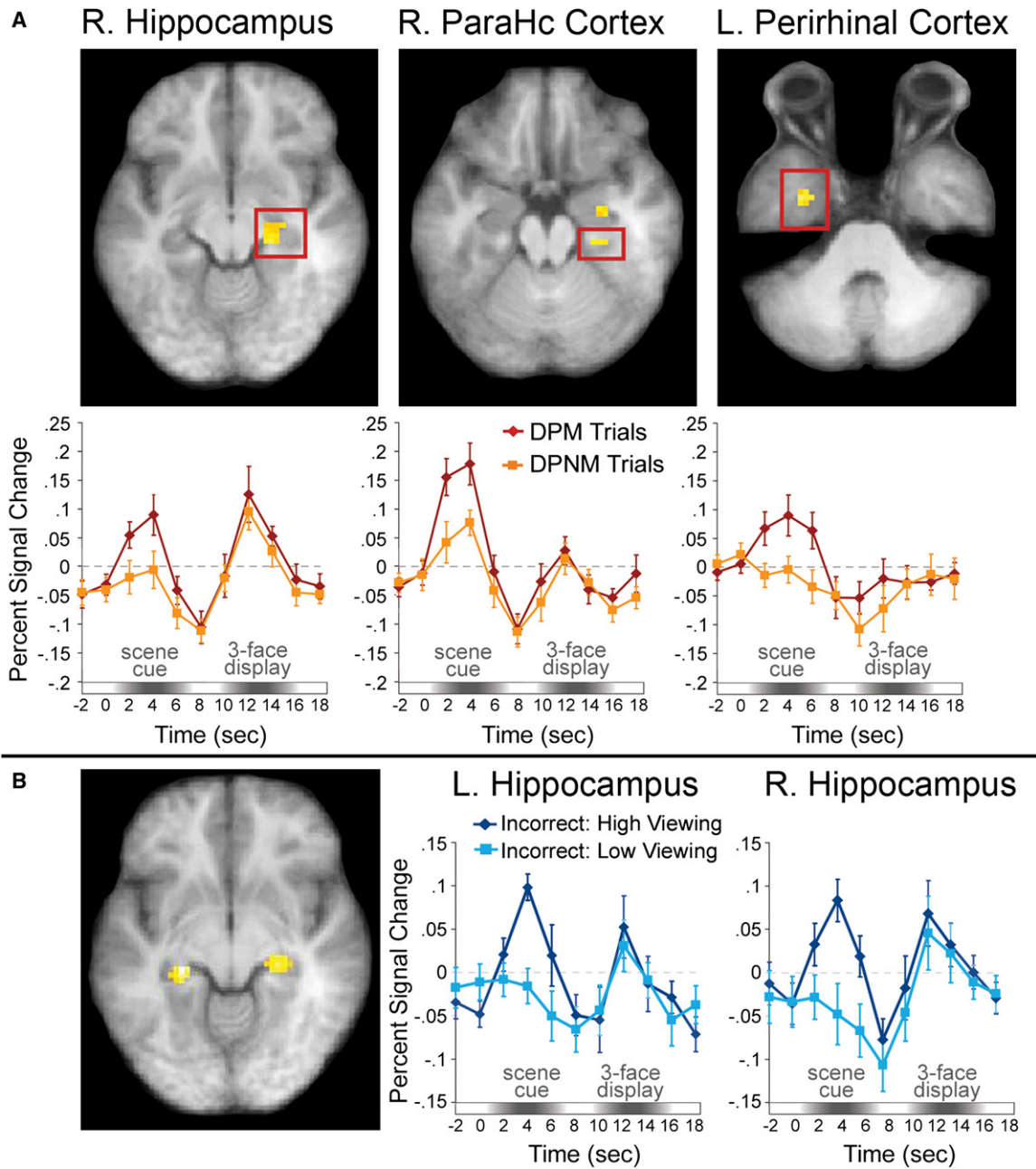


Figure 3. Medial Tempalobe Activity Predicts Eye-Movement-Based Expressions of Relational Memory, Even When Explicit Recognition Has Failed

(A) Examples of MTL regions that showed increased BOLD signal during the scene cue for trials in which participants viewed the matching face disproportionately (DPM trials) versus trials in which they viewed one of the nonmatching faces disproportionately (DPNM trials). Trial-averaged time courses extracted from each ROI illustrate differences in BOLD signal between DPM and DPNM trials during presentation of the scene cue. (B) BOLD signal was greater in both the left and the right hippocampus for incorrect high-viewing trials than for incorrect low-viewing trials. Standard error bars are plotted around the means.

estimates were correlated with estimates of activity in the rest of the brain (Rissman et al., 2004; see Supplemental Data for details). Voxels in the MTL that showed increased correlations with prefrontal ROIs on correct, as compared to incorrect trials were then identified. There were no statistically reliable changes in connectivity between lateral prefrontal regions and MTL

structures during the scene cue or the delay period. During presentation of the three-face test display, however, functional connectivity between the left DLPFC seed region and several hippocampal regions (left anterior hippocampus: $x = -21, y = -18, z = -18; t(13) = 4.01$; left posterior hippocampus: $x = -21, y = -24, z = -9; t(13) = 4.78$; right anterior

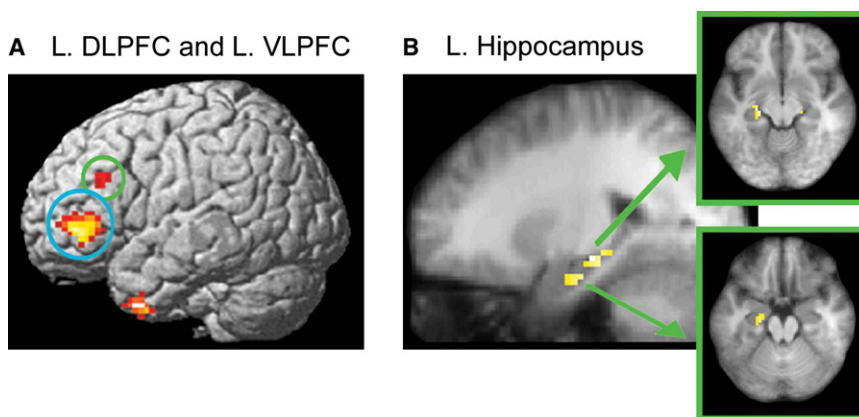


Figure 4. Lateral Prefrontal Activity and Functional Connectivity with the Hippocampus Predicts Accurate Relational Memory Decisions

(A) Regions that showed greater BOLD signal during the scene cue for correct trials than for incorrect trials are rendered on a template brain. Lateral prefrontal areas identified in this contrast are circled (L. DLPFC in green; L. VLPFC in blue). (B) Representative regions in the left hippocampus that exhibited greater connectivity with the left DLPFC seed region on correct than on incorrect trials during presentation of the three-face test display.

hippocampus: $x = 24, y = -21, z = -15; t(13) = 3.53$) was increased on correct, as compared to incorrect trials (see Figure 4B). Functional connectivity was also increased between the left VLPFC seed and regions in the left hippocampus ($x = -21, y = -18, z = -12; t(13) = 3.58$), left parahippocampal cortex ($x = -18, y = -24, z = -21; t(13) = 5.90$), and left perirhinal cortex ($x = -18, y = -6, z = -33; t(13) = 4.36$) during presentation of the three-face test display for correct versus incorrect trials.

DISCUSSION

The aim of the current investigation was to determine whether the hippocampus and adjacent MTL structures support the expression of relational memory in eye-movement behavior, even when behavioral responses are incorrect. Such an outcome would be significant because most theories emphasize the role of MTL structures in conscious retrieval of past events (Aggleton and Brown, 1999; Moscovitch, 1995; Squire et al., 2004; Tulving and Schacter, 1990; Yonelinas, 2002). Results showed that activity in the hippocampus during presentation of the scene cue predicted subsequent viewing of the associated face during the three-face test display, even when participants failed to explicitly identify the match. In contrast, activity in PFC regions was sensitive to subsequent response accuracy but did not predict viewing of matching faces on incorrect trials. Finally, functional connectivity between lateral PFC and hippocampus was increased during presentation of the three-face test display on correct as compared to incorrect trials. Together, these results suggest that hippocampal activity may support the expression of relational memory and that recruitment of a broader network of regions may be required to use this information to guide overt behavior.

Previous evidence taken to support hippocampal contributions to memory without awareness (Chun and Phelps, 1999; Greene et al., 2007; Ryan et al., 2000) has been challenged by recent research (Manns and Squire, 2001; Smith et al., 2006; Preston and Gabrieli, 2008). For example, the failure of amnesic patients to show implicit response facilitation to repeated displays in the contextual cueing task (Chun and Phelps, 1999) has since been attributed to extensive, rather than hippocampally limited, MTL lesions (Manns and Squire, 2001), and results

from a recent fMRI experiment (Preston and Gabrieli, 2008) showed that hippocampal activation during performance of the contextual cueing task was tied to explicit recognition of repeated displays. At first blush, these results may seem to challenge the idea that the hippocampus can support expressions of relational memory without awareness. However, it has been argued that contextual cueing may depend on configural representations supported by extrahippocampal regions such as the perirhinal cortex rather than the kind of relational memory representations thought to depend on the hippocampus (Preston and Gabrieli, 2008).

The results reported here suggest that the hippocampus can support expressions of relational memory even when behavioral responses are incorrect. These results are compelling when considered along with previous findings that show that amnesic patients fail to look disproportionately at relational changes in previously studied scenes even though college-age participants do so despite being unaware of the manipulation (Ryan et al., 2000). Our results also complement previous fMRI research that has shown increased hippocampal activity during presentation of subliminally presented face-occupation pairs (e.g., Degonda et al., 2005) and during implicit learning (Greene et al., 2006; Schendan et al., 2003).

Considered together, these results are consistent with the two-stage model of recollection recently proposed by Moscovitch (2008). According to this model, the initial activation of hippocampal representations (“ecphory”) can guide behavior in an obligatory manner, even before information is consciously apprehended. Thereafter, the individual may become aware of ecphoric output and consciously use this output to guide volitional behavior. The model suggests that hippocampal activity should be correlated with recollection under most circumstances (Eichenbaum et al., 2007) but also suggests that the hippocampus can support expressions of memory even when the second, conscious stage of processing is disrupted.

Although our results indicate that explicit recollection is not a necessary condition for hippocampal recruitment, they do not contradict the idea that hippocampal activity is typically correlated with recollection. Hippocampal activity was not robustly correlated with overt response accuracy in the current experiment, but there are several possible explanations for this null

result (see [Supplemental Data](#) for details). Furthermore, although hippocampal activation has been correlated with recollection in many studies (e.g., [Diana et al., 2009](#)), null results in this area are not uncommon (see [Henson, 2005](#), for review). In general, further work needs to be done to examine the connection between hippocampal activity, eye-movement-based measures of relational memory and explicit recognition accuracy. A full factorial analysis would be needed to address this question, but because viewing of the match was correlated with accuracy, it was not feasible to examine accuracy effects for trials matched on viewing time in the current investigation. Accordingly, an important question for future research is whether hippocampal activity would be greater for correct trials with high viewing of the match than for incorrect trials with high viewing of the match.

As indicated above, activity in the left lateral PFC during processing of the scene cue and functional connectivity between this region and the hippocampus during presentation of the three-face test display was correlated with accurate associative recognition. Previous research implicates left lateral prefrontal regions in retrieval of source information or contextual recollection that may support accurate responses (see [Fletcher and Henson, 2001](#)), and recent work ([Dobbins and Sanghoo, 2006](#)) suggests that left DLPFC in particular may be important for evaluating recovered content with respect to a particular behavioral goal. The present results suggest that the hippocampus may support retrieval of relationally bound information but that regions in the prefrontal cortex may also be recruited to support the use of this information in order to guide explicit associative memory decisions ([Duarte et al., 2005](#)).

The practical implications of the results reported here are potentially far-reaching because they suggest that eye movements provide a powerful approach to investigating relational memory and hippocampal function. Accordingly, eye-tracking may be a valuable tool in translational research, as it is often difficult to overtly assess relational memory in cognitively impaired clinical populations (who may not be able to perform complex meta-cognitive judgments) or in monkeys and rodents (for whom subjective reports of memory retrieval are not possible and must be inferred). Along similar lines, recent work ([Richmond and Nelson, 2009](#)) has demonstrated that this methodological approach is beneficial to memory studies conducted with infants, who cannot yet report the contents of what has been successfully retrieved from memory. Finally, eye-tracking could be used to obtain information about past events from participants who are unaware or attempting to withhold that information. In other words, there may be circumstances in which eye movements provide a more veridical and robust account of past events or experiences than behavioral reports alone.

EXPERIMENTAL PROCEDURES

Participants

Participants were 18 right-handed individuals (eight women) from the UC Davis community who were paid in exchange for participation. Four participants were excluded from the reported analyses; one because behavioral performance was at chance and the remaining three because eye position could not be reliably monitored.

Procedure

After informed consent was obtained and instructions were provided, each participant practiced the face-scene association task (see below). Scanning commenced once the experimenter was satisfied with the participant's comprehension of the task. The scanning session consisted of four study blocks, each followed immediately by a corresponding test block. Eye position was monitored throughout the entire scanning session, and the eye-tracker was calibrated using a 3 × 3 spatial array prior to the initiation of each experimental block (example stimuli are illustrated in [Figure 1](#) and more detailed information about stimuli and counterbalancing can be found in the [Supplemental Data](#)).

Each study block consisted of 54 study trials, in which a unique scene was presented for 1 s, after which a single face was superimposed on top of that scene for 2 s. To elicit reasonably high levels of accuracy, participants were instructed to determine whether each person denoted by the face looked like they belonged in the place depicted in the scene. A variable duration inter-trial interval (ITI; range 1–5 s) separated subsequent trials, and a white fixation cross was presented centrally during the final 500 ms of the ITI to warn participants that the next trial was about to begin (see [Figure 1](#)). Participants were told that they should orient their gaze to this fixation cross in preparation for the next trial, but that they could move their eyes freely once the scene was presented. The total duration of each study block (i.e., scanning run) was 336 s, including a 12 s unfilled interval at the beginning of each block.

Each test trial (18 per block) began with the presentation of a scene that had been viewed in the previous study block ("scene cue"). The scene remained on the screen for 1 s and was followed by a 7 s delay. Participants were instructed that they should use the scene as a cue to retrieve the associated face before the three-face test display was presented. A white fixation cross, presented in the center of the screen during the final 500 ms of the delay period, encouraged participants to orient their gaze toward the center of the screen in anticipation of the three-face test display, which remained on the screen for 2 s. When the test display was presented, participants were to indicate, via button press, which face (left, right, or bottom) had been paired with that scene earlier. Participants were also given the option to respond "don't know" if they were unsure about the identity of the match and speed was emphasized, but not at the expense of accuracy. A variable duration ITI (range 10–14 s) separated subsequent trials, and a centrally located white fixation cross, presented in the final 500 ms of the ITI, warned participants that the next trial was about to begin (see [Figure 1](#)). The total duration of each test block (i.e., scanning run) was 408 s, including a 12 s unfilled interval at the beginning of each block.

Eye-Tracking Acquisition and Analysis

Eye position was monitored during fMRI scanning at a rate of 60 Hz using an MRI-compatible Applied Science Laboratories (ASL) 504 long-range optics eye tracker. Eye-tracking analyses focused on eye movements that occurred during the 2 s following three-face display onset. Fixations made during this period were assigned to particular regions of interest (ROIs) within each three-face test display (i.e., left face, right face, bottom face, background scene), and the proportion of total viewing time allocated to each ROI was calculated (see [Supplemental Data](#) for details).

Image Acquisition and Preprocessing

fMRI data were acquired with a 3T Siemens Trio scanner located at the UC Davis Imaging Research Center. Functional data were obtained with a gradient echoplanar imaging (EPI) sequence (repetition time, 2000 ms; echo time, 25 ms; field of view, 220; 64 × 64 matrix); each volume consisted of 34 axial slices, each with a slice thickness of 3.4 mm, resulting in a voxel size of 3.4375 × 3.4375 × 3.4 mm. Coplanar and high-resolution T1-weighted anatomical images were acquired from each participant, and a simple motor-response task ([Aguirre et al., 1998](#)) was performed to estimate subject-specific hemodynamic response functions (HRF).

Preprocessing was performed using Statistical Parametric Mapping (SPM5) software. EPI data were slice-timing corrected using sinc interpolation to account for timing differences in acquisition of adjacent slices, realigned using a six-parameter, rigid-body transformation, spatially normalized to the Montreal Neurological Institute (MNI) EPI template, resliced into 3 mm isotropic

voxels, and spatially smoothed with an isotropic 8 mm full-width at half-maximum Gaussian filter.

fMRI Data Analysis

Event-related BOLD responses associated with each component of each test trial (i.e., scene cue, delay, and three-face test display) were deconvolved using linear regression (Zarahn et al., 1997). Covariates of interest were created by convolving vectors of neural activity for each trial component with subject-specific HRFs derived from responses in the central sulcus for each participant during a visuomotor response task. Data from the visuomotor response task were unavailable or unreliable for four participants. For these individuals, covariates were constructed by convolving neural activity vectors with an average of empirically-derived HRFs from 18 participants.

Separate covariates were constructed to model responses for each test trial component (scene cue, delay, three-face display) as a function of viewing time (i.e., DPM versus DPNM trials), viewing time for incorrect trials only (i.e., incorrect trials with high viewing of the match versus incorrect trials with low viewing of the match), and behavioral response accuracy (i.e., correct versus incorrect identification of the matching face). Each classification scheme resulted in six distinct covariates of interest that modeled activity during each task phase either as a function of eye-movement-based memory measures (scene cue – disproportionate match, scene cue – disproportionate nonmatch, etc.), eye-movement-based memory measures for incorrect trials (scene cue – incorrect high viewing, scene cue – incorrect low viewing, etc.), or response accuracy (scene cue – correct, scene cue – incorrect, etc.). Additional covariates of no interest modeled spikes in the time series, global signal changes that could not be attributed to variables in the design matrix (Desjardins et al., 2001), scan-specific baseline shifts, and an intercept. Regression analyses were then performed on single-subject data using the general linear model with filters applied to remove frequencies above 0.25 Hz and below 0.005 Hz. These analyses yielded a set of parameter estimates for each participant, the magnitude of which can be interpreted as an estimate of the BOLD response amplitude associated with a particular trial component (e.g., responses during the scene cue on DPM trials).

After single-subject analyses were completed, images for the contrasts of interest were created for each participant. Contrast images were entered into a second-level, one-sample *t* test, in which the mean value across participants for every voxel was tested against zero. Significant regions of activation in the MTL were identified using an uncorrected threshold of $p < 0.005$ and a minimum cluster size of eight contiguous voxels. With this voxel-wise threshold, the family-wise error rate for the MTL (i.e., hippocampus, parahippocampal, perirhinal, and entorhinal cortices), estimated using a Monte Carlo procedure (as implemented in the AlphaSim program in the AFNI software package), was constrained at $p < 0.05$. Because we predicted that relational memory retrieval would be triggered by presentation of the scene cue, analyses reported here focus on this task period.

Detailed information about the number of trials per bin for each participant in every contrast is provided in Table S4. Because bin sizes for incorrect viewing time analyses were small for some participants, additional analyses were conducted to examine the reliability of incorrect high- versus low-viewing time effects in the fMRI data when these participants were excluded. Results of these analyses are consistent with those reported in the manuscript (see Table S5).

Additional analyses were performed to identify regions outside of the MTL for which activity during presentation of the scene cue was correlated with eye-movement behavior and response accuracy. These regions were identified using an uncorrected threshold of $p < 0.001$ and a minimum cluster size of 8 contiguous voxels. Coordinates of local maxima from these contrasts during presentation of the scene cue are summarized in Tables S1–S3.

SUPPLEMENTAL DATA

Supplemental Data include Supplemental Experimental Procedures, Results, Discussion, five figures, and five tables and can be found with this article online at [http://www.cell.com/neuron/supplemental/S0896-6273\(09\)00636-9](http://www.cell.com/neuron/supplemental/S0896-6273(09)00636-9).

ACKNOWLEDGMENTS

This work was supported by a National Institute of Mental Health Fellowship F32MH075513 (D.E.H.) and a National Institutes of Health Grant MH68721 (C.R.). The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institute of Mental Health or the National Institutes of Health. We are very grateful to Dr. Neal Cohen for insightful comments and for sharing the materials that were used in this experiment. Many thanks as well to Charlie Newer and Evan Katsurani for technical guidance and assistance with data collection.

Accepted: August 21, 2009

Published: September 9, 2009

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