

UC Davis

UC Davis Previously Published Works

Title

The Hippocampus Generalizes across Memories that Share Item and Context Information

Permalink

<https://escholarship.org/uc/item/77b169zg>

Journal

Journal of Cognitive Neuroscience, 31(1)

ISSN

0898-929X

Authors

Libby, Laura A

Reagh, Zachariah M

Bouffard, Nichole R

et al.

Publication Date

2019

DOI

10.1162/jocn\_a\_01345

Peer reviewed



Published in final edited form as:

*J Cogn Neurosci*. 2019 January ; 31(1): 24–35. doi:10.1162/jocn\_a\_01345.

## The hippocampus generalizes across memories that share item and context information

Laura A. Libby<sup>1</sup>, Zachariah M. Reagh<sup>1,2</sup>, Nichole R. Bouffard<sup>1</sup>, J. Daniel Ragland<sup>3</sup>, Charan Ranganath<sup>1,4</sup>

<sup>1</sup>Center for Neuroscience, University of California, Davis

<sup>2</sup>Department of Neurology, University of California, Davis

<sup>3</sup>Department of Psychiatry and Behavioral Sciences, University of California, Davis

<sup>4</sup>Department of Psychology, University of California, Davis

### Abstract

Episodic memory is known to rely on the hippocampus, but how the hippocampus organizes different episodes to permit their subsequent retrieval remains controversial. One major area of debate hinges on a discrepancy between two hypothesized roles of the hippocampus: differentiating between similar events to reduce interference, and assigning similar representations to events that share overlapping items and contextual information. Here, we used multivariate analyses of activity patterns measured with functional magnetic resonance imaging (fMRI) to characterize how the hippocampus distinguishes between memories based on similarity at the level of items and/or context. Hippocampal activity patterns discriminated between events that shared either item or context information, but generalized across events that shared similar item-context associations. The current findings provide evidence that, whereas the hippocampus can reduce mnemonic interference by separating events that generalize along a single attribute dimension, overlapping hippocampal codes may support memory for events with overlapping item-context relations. This lends new insights into the way the hippocampus may balance multiple mnemonic operations in adaptively guiding behavior.

### INTRODUCTION

Memories for past events include information about who or what was encountered (“items”), as well as information about the place, time, and situation in which the event took place (“context”). For instance, seeing a friend’s dog (an item) at the neighborhood park (a context) might remind you of the time you played with that dog in your friend’s backyard (same item, different context), or you might remember that, the last time you visited the park, you ran into a coworker (different item, same context). This example illustrates how the human brain can both distinguish between specific events and generalize across events according to similarity in item and/or context information (Johnson et al., 1993, Mitchell and Johnson, 2009). Memory for item-context associations is known to rely on the

hippocampus (Cohen et al., 1997, Vargha-Khadem et al., 1997, Brown and Aggleton, 2001, Scoville and Milner, 1957). However, it is unclear how the hippocampus organizes memories to facilitate item and context retrieval when content is highly overlapping, as in many of our real world experiences.

One prominent model holds that the hippocampus has a computational specialization for mapping distinct events onto orthogonal neural codes, a process known as pattern separation (Norman and O'Reilly, 2003, Marr, 1971, Rolls and Kesner, 2006, Yassa and Stark, 2011). Pattern separation is thought to prevent catastrophic interference between representations with similar features; when hippocampal pattern separation fails, episodic memory is predicted to break down due to overgeneralization (Norman and O'Reilly, 2003, Norman, 2010). Results from single-unit recording studies in rats support this view, showing that spatial coding in the hippocampus dramatically “remaps” with slight changes in spatial context (Guzowski et al., 2004, Leutgeb et al., 2007, Bostock et al., 1991, Lever et al., 2002, Neunuebel and Knierim, 2014, Wills et al., 2005). Studies in rodents have not yet characterized a role for the hippocampus in pattern separation of items independent of their spatial context. However, several human neuroimaging studies measuring the similarity of hippocampal voxel activity patterns across trials as an indirect measure of neuronal population coding (Kriegeskorte et al., 2006, Kriegeskorte et al., 2008) have found that hippocampal activity patterns tend to differentiate between similar items to a greater extent than cortical regions (Berron et al., 2016, Huffman and Stark, 2014, Liang et al., 2012). Further evidence (LaRocque et al., 2013) suggests that greater hippocampal differentiation between similar items during learning predicts memory for these items after a delay.

A second framework for understanding hippocampal function suggests that, whereas item and context information are respectively processed in distinct neocortical pathways through perirhinal and parahippocampal cortex (Ranganath and Ritchey, 2012, Ritchey et al., 2015a), the hippocampus specifically represents item-context associations (Eacott and Gaffan, 2005, Davachi, 2006, Ranganath, 2010, Knierim et al., 2006). Consistent with this idea, single-unit recording studies in rats have shown that hippocampal neuronal ensemble coding for items encountered in a particular spatial context is highly reliable across repeated exposures (McKenzie et al., 2014, Manns and Eichenbaum, 2009). Complementary results in humans have shown that hippocampal voxel patterns are similar when the same item is presented repeatedly in the same position within a temporal context (Hsieh et al., 2014, Libby et al., 2014, Ritchey et al., 2015b). There is also evidence for similar item-level voxel patterns in the human hippocampus based on both spatial and temporal proximity in virtual environments (Deuker et al., 2016). Finally, hippocampal activity patterns carry information about item-context bindings in the case of even imagined links between objects and spatial locations (Sheldon and Levine, 2015).

The “pattern separation” (Norman and O'Reilly, 2003, Marr, 1971, Rolls and Kesner, 2006, Yassa and Stark, 2011) and “items-in-context” (Eacott and Gaffan, 2005, Davachi, 2006, Ranganath, 2010, Knierim et al., 2006) views are largely complementary, in that the former addresses how the hippocampus organizes distinct events with respect to each other and the latter focuses on the relative importance of item-context associations in episodic memory. However, a fundamental and currently unresolved question is how these processes coexist

and interact. A strong version of the view emphasizing pattern separation would suggest that the hippocampus sharply differentiates between events, even if they share information along one or more attribute dimensions (i.e., item and/or context information). That is, the tendency to orthogonalize similar inputs may override associative binding. A strong version of the items-in-context perspective, in turn, would suggest that the hippocampus assigns overlapping neural representations to events in which similar items were encountered in similar contexts. That is, context stability may override the demand to orthogonalize similar inputs. The stability and generalization of hippocampal codes may also map onto different regions along its longitudinal axis (Poppenk et al., 2013), as the anterior hippocampus has been hypothesized to support coarse, generalized representations compared to more granular representations in the posterior extent (Brunec et al., 2018).

To address these outstanding questions, we used functional magnetic resonance imaging (fMRI) to examine hippocampal activity patterns during a memory test that required retrieval of both item and context information. Critically, we manipulated overlap of both item and/or context information: Different exemplars from the same item category were learned in each of four different context locations, resulting in sets of events with overlapping item attributes, context associations, or both (Figure 1A). Participants were then scanned during a recognition test that required recall of item-context relations (Figure 1B). We analyzed the similarity of voxel activity patterns evoked during successful retrieval of item-context relations to estimate the extent to which brain areas differentiated or generalized across memories with overlapping item, context, and item-in-context information (Figure 2A). In addition to whole-brain activity patterns, we focused specifically on the hippocampus (including head/body/tail subdivisions), as well as perirhinal and parahippocampal cortex for comparison.

## MATERIALS AND METHODS

### Participants

Twenty-five healthy human participants (female=11; mean age=24.4 years; standard deviation=4.6 years) underwent fMRI scanning during a long-term memory encoding and retrieval task. Participants with head motion greater than 3 mm from origin (N=2) or chance-level context memory performance (N=3) were excluded from analysis, resulting in a total of 20 included participants (female=8). All procedures were approved by the University of California, Davis Institutional Review Board.

### Materials

Item stimuli consisted of 336 visual objects evenly distributed across 28 categories (e.g., toasters, dogs, jackets, muffins), with 12 unique exemplars per category. For each participant, within each category, eight objects were randomly assigned to the list of target stimuli and four objects were assigned to the list of foils, for a total of 224 targets (presented in the encoding and retrieval phases) and 112 foils (presented only in the retrieval phase). Target stimuli were then randomly assigned to one of four encoding context locations (either park, beach, restaurant, or café), for a total of 56 stimuli per context. Two exemplars from each category were presented in each context.

## Procedure

**Context personalization and task practice phase – Behavioral**—During the encoding phase, participants were cued to visualize each item in one of four spatial contexts: a park, a beach, a restaurant, or a café; at test, participants were asked to retrieve the context originally associated with each item. To ensure that the locations evoked by context cues were visualizable and consistent over the course of the task, context cues were personalized for each participant prior to the start of the task. Participants were asked to select a specific park, beach, restaurant, and café from their past experience that they could picture vividly, and were then instructed to bring only those venues to mind in response to the context cues. Participants then received task instructions and completed practice encoding and retrieval runs (using the same four context cues but an independent practice set of item stimuli) on a computer outside of the scanner. Participants were permitted to repeat the practice runs as many times as was necessary to become comfortable with the task timing and response mappings.

## Encoding phase

Over four encoding runs (Figure 1A), target items were presented sequentially for 2 s each. Item trials were broken down into mini-blocks varying in size from two to five items per block. At the beginning of each mini-block, one of four context cue screens was presented for 3 s, either “PARK,” “BEACH,” “RESTAURANT,” or “CAFÉ,” cuing participants to visualize a specific context location for the duration of the mini-block. Then, for each item in the mini-block, participants visualized the item in that particular context location and made a yes/no response (via a button press) to the question, “In real life, would you be likely to see this item in this location?” Item category, context location, and item category-context pairings were counterbalanced across runs. Within run, item order was pseudo-randomized for each participant, with the constraint that no adjacent item trials contained exemplars from the same category. Block order was also pseudo-randomized for each participant such that no adjacent mini-blocks contained the same context cue, and block length was counterbalanced across context question and across scanning run. Context cue and item trial screens were presented with a jittered ISI optimized for event-related fMRI (SOA 4–12 s with a mean of 6 s).

## Retrieval phase

Immediately following each encoding run, participants completed a mixed item recognition/source memory test during fMRI scanning (Figure 1B). All target items from the most recent encoding run and a full set of category-matched foils were presented sequentially for 2 s each. For each trial, participants indicated the quality and content of their memory for that item by selecting one of seven response button options: 1) the item was novel (“New”); 2) they were familiar with the item but could not recollect any contextual details associated with having seen it before (“Familiar”); 3) they recollected having seen the item before and could bring some contextual information to mind but *not* the specific location with which the item was originally associated (“Recollect”); 4–7) if they recollected the item and could remember which of the four encoding context locations was originally associated with that item, they pressed a button indicating that location (“Park,” “Beach,” “Restaurant,” or

“Café”). Retrieval trials were pseudo-randomized such that no adjacent trials contained exemplars from the same category and jittered for event-related fMRI (SOA=4–12 s with a mean of 6 s). In total, four encoding runs were interleaved with four retrieval runs.

### Image acquisition

MRI scanning was conducted at the UC Davis Facility for Integrative Neuroscience on a 3T Siemens Skyra with a 32-channel phased-array head coil. High-resolution T1-weighted structural images were acquired using a magnetization-prepared rapid acquisition gradient echo (MPRAGE) pulse sequence (1 mm<sup>3</sup> voxels; matrix size=256 × 256; 208 slices). Images sensitive to BOLD contrast were acquired using a whole-brain multiband gradient echo planar imaging (EPI) sequence (3 mm<sup>3</sup> voxels; TR=1220 ms; TE=24 ms; FA=67°; multiband acceleration factor=2; 38 interleaved slices; FOV=192 mm; matrix size=64 × 64) during task performance.

### Behavioral analysis

For the purposes of the fMRI-based multivoxel pattern similarity analysis, detailed below, it was important to establish: 1) that correct context memory judgments were likely to be driven by real memory signal, and 2) whether there were any systematic differences in item recognition or context memory across encoding contexts. Discriminability (via the  $d'$  sensitivity index) was calculated as the z-scored hit rate minus the z-scored false alarm rate. For item discriminability, hit rate was calculated as the percentage of target trials given any “old” item response (“Familiar,” “Recollect,” or any of the encoding context location judgments – even if the specific location was incorrect) for items encoded in each context location – and overall item false alarm rate was calculated as the proportion of “old” item responses given on foil trials. Differences in item  $d'$  across encoding contexts were tested via a one-way repeated measures ANOVA (with a Huynh-Feldt correction for non-sphericity).

To estimate the meaningfulness of context memory judgments, context discriminability ( $d'$ ) was calculated separately for each encoding context as the difference between the probability of a correct context judgment (e.g., an item encoded in the café given a “Café” response at test) and the probability of a false alarm to that particular context location (e.g., an item encoded in the park or a foil item given a “Café” response at test). Subjects with context  $d'$  at floor (i.e., not significantly above chance) for any encoding context were excluded from fMRI analysis (N=3). Context discriminability estimates were entered into a one-way repeated measures ANOVA to test for differences across encoding contexts (with a Huynh-Feldt correction for non-sphericity).

### Multivoxel pattern similarity analysis

Because we were interested in understanding how the MTL organizes item and context information during successful reinstatement of item-context relations, multivoxel pattern similarity analysis focused on fMRI data from the retrieval phase. EPI timeseries underwent motion correction and highpass filtering (0.01 Hz) in FMRIB’s Software Library (FSL). In preparation for multivoxel pattern similarity analysis procedures, spatial smoothing was omitted. Event-related blood oxygenation level-dependent (BOLD) signal change was

estimated separately for each item trial, controlling for signal change due to all other trials and motion artifact, using ordinary least squares regression, resulting 336 single-trial beta images (Xue et al., 2010, Mumford et al., 2012). Single-trial beta images from runs 2–4 were coregistered with single-trial beta images from run 1 using FSL’s FLIRT linear registration software (6 degrees of freedom). Coregistered single-trial beta images with atypically high mean absolute z-score (based on the distribution of beta estimates for each grey matter voxel across all trials) were excluded from further analysis (mean absolute z threshold = 1.5; between 0 and 10 trials excluded per subject, median = 4.5 trials). This objective noise trial exclusion procedure has been effective in previous pattern similarity studies (Libby et al., 2014), and, in the current study, beta images were additionally visually inspected to verify that subjectively noisy trials were not overlooked. Critically, to pinpoint retrieval trials containing both (perceived) item and (reinstated) context information, the subsequent pattern similarity analysis was restricted to correct context memory judgment trials.

To determine the spatial distribution of information coding effects across the hippocampus and surrounding cortical areas, we employed multivoxel pattern similarity analysis (Kriegeskorte, 2011, Kriegeskorte et al., 2008) using a searchlight approach (Kriegeskorte et al., 2006). For every voxel in the brain, correlation coefficients (Pearson’s  $r$ ) were calculated for all pairs of trials based on the pattern of beta coefficients contained in a sphere with a 5-voxel diameter centered on that voxel, resulting in an observed pattern similarity (correlation) matrix. Searchlight spheres centered within a given ROI excluded voxels from other ROIs. To reduce the influence of temporal autocorrelation and within-run dependencies on pattern information coding effects (Mumford et al., 2014), observed similarity estimates from trial pairs within the same fMRI run were discarded. Between-run observed pattern similarity estimates were entered into second-order similarity analyses. Here, we refer to “second-order similarity” as the z-transformed point-biserial correlation between observed pattern similarity estimates and an “ideal” binary model of predicted pattern similarity (Figure 2B), where trial pairs that were similar along a dimension of interest were coded as 1 and trial pairs that differed along that dimension were coded as 0. In this case, the point-biserial correlation is effectively identical to a two-sample t-test, and was selected because it is sensitive to within-subject variance, providing a somewhat conservative metric of multivoxel pattern similarity.

We examined pattern similarity effects based on shared item and context information in two separate analyses with orthogonal predicted pattern similarity models. To identify brain regions where voxel patterns carried information about the item category, a binary model was constructed labeling trial pairs according to whether they contained items from the same category (e.g., two dog trials – “*similar item*” pairs) or different categories (e.g., one dog trial and one toaster trial – “*different item*” pairs), excluding trial pairs with the same context location. To identify brain regions with voxel patterns carrying retrieved context information, a binary model was constructed labeling trial pairs according to whether the retrieved context was the same (e.g., two park trials – “*similar context*” pairs) or different (e.g., a park trial and a beach trial – “*different context*” pairs), excluding trial pairs with the same item category. The reason we here refer to shared contexts as “similar” is because although the context is ostensibly the same, we cannot be sure that its mental construction

identical across experiences. Moreover, it is likely that different items have unique associations with the same context, leading to subtle differences in spite of overall stability.

To test for pattern similarity effects driven by shared item-context relations that were distinguishable from item or context effects alone, we constructed a binary model for item-in-context similarity that was superordinate to the item and context models. Trial pairs with both item category and context location in common (e.g., two dog-in-the-park trials – “*similar item+context*” pairs) were coded as 1 and trial pairs with overlapping item category *or* context location, but not both (i.e. *similar item* and *similar context* pairs), were coded as 0; trial pairs that were different on both item category and context location dimensions were excluded. This model, as well as the similar item and similar context models are visualized in Figure 2B.

For each of the three predicted similarity models, the resulting second-order similarity estimate was assigned to the center voxel of each searchlight, resulting in three whole-brain pattern similarity images for each subject. Single-subject pattern similarity images were normalized to the MNI152 template via the application of linear transformation matrices calculated in a two-step registration process: using FLIRT, representative single-subject EPI images were co-registered with high-resolution anatomical MPRAGE images (6 degrees of freedom), MPRAGE images were normalized to the template image (12 degrees of freedom), and the two resulting transformation matrices were mathematically concatenated (multiplied). Brain regions that reliably carried each type of information across subjects ( $p_{\text{FWE}} < 0.05$ ) were identified non-parametrically based on the distribution of maximum cluster mass (after a voxelwise t-threshold of 2.539) across 10,000 permutations of the data (sign-flipping approach) using the Randomise function in FSL (Nichols and Holmes, 2002). Because we had *a priori* interest in the hippocampus and surrounding cortex, group analyses were restricted to MTL voxels using a statistically conservative (anatomically liberal) MTL mask generated with the WFU Pickatlas (Maldjian et al., 2003) consisting of parahippocampal gyrus, hippocampus, uncus, and amygdala and dilated in three dimensions by 4 mm. The dilated anatomical MTL mask extended laterally and ventrally into fusiform cortex, posteriorly to the level of the posterior horn of the lateral ventricles, and anteriorly into medial temporopolar cortex. We additionally explored pattern similarity effects outside of the MTL using the same non-parametric permutation test procedure, but with whole-brain grey matter masks ( $p(\text{grey matter}) > 0.36$ , corresponding to a threshold of 90 out of 245 of the FSL MNI 152 segmentation grey matter priors) that excluded voxels contained in the MTL mask.

To allow for visualization and comparison of pattern similarity effects specifically in the hippocampus, for each subject, a second set of searchlight images was generated containing average pattern similarity estimates for each combination of item category and context location overlap. That is, one image contained average similarity estimates for *similar item* pairs, one image contained average similarity estimates for *similar context* pairs, one image contained the average across *similar item+context* pairs, and one image contained the average across trial pairs that were different on both item category and context location (“*both different*” pairs). From these images, mean pattern similarity estimates were extracted using individually-defined regions of interest (ROIs) corresponding to head, body, and tail



divisions of the hippocampus. Because items-in-context and pattern separation accounts of MTL function both predict that adjacent MTL cortical areas tend to generalize across similar events, pattern similarity estimates were also extracted from anatomical ROIs in perirhinal cortex and parahippocampal cortex. All ROIs were defined separately for right and left hemispheres, and anatomical landmarks for ROI definition are thoroughly described by Moore et al. (2014).

Pattern similarity estimates were entered into a full factorial repeated measures ANOVA with ROI and information overlap as factors of interest (using a Huynh-Feldt correction for non-sphericity), controlling for hemisphere. Significant interactions were broken down separately by ROI. *A priori* contrasts testing for item generalization (*similar item* > *both different*), context generalization (*similar context* > *both different*), and item-in-context generalization (*similar item+context* > *similar item* + *similar context*) were defined for information overlap, in keeping with the prediction model hierarchy described above. When two or more *a priori* contrasts were significant within an ROI, we also applied *post hoc* contrasts (with Bonferroni correction for multiple comparisons) to determine which significant effect was strongest (e.g., *similar item+context* > *similar item*). Contrasts were additionally applied to pattern similarity estimates extracted from eroded hippocampal head ROIs that excluded edge voxels (binary erosion in three dimensions with 3mm spherical kernel) to reduce any possible influence of signal from neighboring voxels.

## RESULTS

Because we were interested in voxel patterns associated with memory for items in context, it was important to establish the presence of two behavioral effects: 1) that memory (particularly context memory) *existed*, i.e. that any given correct context memory response could be attributed to real memory signal; and 2) that there were no systematic differences in memory performance across encoding contexts. One-way repeated measures ANOVAs confirmed that both item discriminability ( $F_{(1,19)} = 238.34, p < 0.0001$ ) and context discriminability ( $F_{(1,19)} = 77.20, p < 0.0001$ ) were significantly different from zero across encoding contexts, but that neither metric differed significantly between encoding contexts ( $F$ 's < 1.39,  $p$ 's > 0.25) (Figure 3).

To examine the neural organization of memories for similar events, we performed a set of multivoxel pattern similarity searchlight analyses (Kriegeskorte et al., 2006, Kriegeskorte et al., 2008). Analysis was restricted to retrieval phase trials characterized by accurate context recall based on an item cue. Local voxelwise patterns of evoked activity were correlated between all pairs of these trials, and trial pair correlations were contrasted according to the similarity of item and/or context information contained in each pair (Figure 2).

We first investigated whether local voxel patterns in the hippocampus generalized across different exemplars of the same item. Voxelwise pattern similarity was contrasted between trial pairs containing different item exemplars from the same category ("*similar item*" pairs) against pattern similarity between trial pairs that included exemplars from different item categories ("*both different*" pairs). To isolate item-level pattern information, this analysis excluded trial pairs that had been associated with the same encoding context. In the medial

temporal lobe (MTL), significant clusters extended throughout the ventral and medial temporal neocortex, including perirhinal cortex, and amygdala, but no hippocampal voxels were identified as generalizing across events on the basis of item information alone (Figure 4A), consistent with previous studies of item pattern separation in the hippocampus and medial temporal cortex (LaRocque et al., 2013, Huffman and Stark, 2014, Liang et al., 2012). Across the whole brain, item-level pattern information was identified extensively in frontal, parietal, and occipital cortex, with the strongest effects in right lateral occipital cortex and posterior fusiform cortex (Figure 4A).

We next examined whether the hippocampus generalized across different items according to whether they had been studied with the same encoding context. We contrasted voxel pattern similarity between trial pairs with different item cues that shared a study context (“*similar context*” pairs) and *both different* pairs. No suprathreshold clusters in hippocampus or neocortex were identified as generalizing across events on the basis of contextual information, either within the MTL or across the whole brain. Given that items in this study have concrete sensory attributes whereas contexts are imagined, subtler context-driven effects are perhaps not surprising. To enhance our sensitivity to detect effects of shared context, we examined whether the strength of neural pattern similarity evidence for context reinstatement might be related to individual differences in behavioral evidence for context memory. In a voxelwise regression (with non-parametric inference testing, as above, but with full permutations), we identified regions where pattern similarity evidence related to context similarity were positively associated with individual behavioral context discriminability estimates (averaged across contexts). Family-wise error rate was determined on the basis of cluster mass using independent MTL and non-MTL grey matter masks. Because no clusters in the MTL surpassed the stringent cluster mass-based family-wise error rate correction applied in other analysis, we additionally applied threshold-free cluster enhancement (TFCE) to detect the presence of any relatively weaker but reliable voxelwise effects ( $p < 0.05$ , FWE-corrected) (Smith & Nichols, 2009). Pattern similarity driven by retrieval of similar contexts was positively associated with context discriminability across subjects in parahippocampal cortex, retrosplenial cortex, lingual gyrus, and precuneus (Figure 4B). However, again, no significant hippocampal voxels were identified.

Our third analysis tested the possibility that the hippocampus generalized across memories that included similar item and context information. If the hippocampus organizes memories on the basis of item-context associations, voxel pattern similarity should be substantially higher across events with similar items and similar contexts, compared to events that shared only a single attribute. Therefore, we contrasted voxel pattern similarity between trial pairs with that included similar items that had been associated with the same study context location (“*similar item+context*” pairs) against pattern similarity between trial pairs that overlapped on one, but not both, of these dimensions (i.e., *similar item* and *similar context* pairs). Searchlight analysis identified clusters showing this effect with centers of mass in the anterior and middle hippocampus bilaterally, extending partially into ventral and medial temporal neocortex (Figure 4C).

To further characterize item and context organization within the hippocampus and adjacent cortical regions, mean pattern similarity estimates (z-scores) for *both different*, *similar item*,

*similar context*, and *similar item+context* pairs were extracted from anatomically-defined hippocampal and cortical regions of interest (ROIs). Because of previously reported functional differences along the anterior-posterior axes of the hippocampus (Poppenk et al., 2013), we evaluated hippocampal head, body, and tail separately using established anatomical landmarks (Moore et al., 2014). A full factorial repeated measures ANOVA revealed a significant main effect of ROI ( $F_{(4,19)}=16.37$ , corrected  $p_{HF} < 0.0001$ ,  $\epsilon_{HF}=0.54$ ), a significant main effect of information overlap ( $F_{(3,19)}=4.45$ , corrected  $p_{HF} < 0.05$ ,  $\epsilon_{HF}=0.41$ ), and, critically, a significant ROI-by-information overlap interaction ( $F_{(12,19)}=4.25$ , corrected  $p_{HF} < 0.005$ ,  $\epsilon_{HF}=0.36$ ); there was no significant main effect of hemisphere or interaction between hemisphere and either factor of interest ( $F$ 's  $< 1.58$ ,  $p$ 's  $> 0.18$ ). Based on the significant interaction, we examined the effect of information overlap within each ROI, controlling for hemisphere, using a set of *a priori* and follow-up *post hoc* contrasts to detect the presence of item, context, and item-in-context generalization separately for each ROI.

In addition to the hippocampus, we examined the extent to which MTL cortical ROIs generalized across items as points of comparison. Within perirhinal cortex, there was a trend toward greater pattern similarity estimates for *similar item* compared to *both different* trial pairs ( $F_{(1,19)}=3.12$ ,  $p=0.08$ ), and similarity estimates were greater for *similar item+context* relative to *similar item* pairs ( $F_{(1,19)}=12.9$ ,  $p < 0.005$ , Bonferroni-corrected), suggesting that item generalization in perirhinal cortex is strengthened by context similarity (Figure 5A). In parahippocampal cortex, pattern similarity between *similar item* pairs was significantly greater than *both different* pairs ( $F_{(1,19)}=8.38$ ,  $p < 0.01$ ), but were not significantly different from *similar item+context* pairs ( $F_{(1,19)}=12.9$ ,  $p < 0.005$ , Bonferroni-corrected), suggesting that item generalization in parahippocampal cortex was insensitive to context similarity (Figure 5B).

In contrast to item generalization effects in the cortex, within hippocampal head, pattern similarity estimates were significantly and selectively greater for *similar item+context* pairs compared to *similar item* and *similar context* pairs ( $F_{(1,19)}=5.19$ ,  $p < 0.05$ ), consistent with voxelwise results (Figure 5C). This pattern of results suggests that hippocampal head only generalized across trials that shared similar item-in-context information. Eroded hippocampal head ROIs excluding edge voxels demonstrated a highly similar pattern of results, although the selective *similar item+context* effect was attenuated somewhat ( $F_{(1,19)}=3.68$ ,  $p=0.06$ ). Moreover, we note the overlap between these ROI-based results and the whole-brain searchlight results from the prior section. This suggests that the generalization effect in the hippocampal head was not driven by edge voxels, which could potentially include signal outside the hippocampus. There were no significant pattern similarity differences in hippocampal body or tail ROIs ( $F$ 's  $< 1.72$ ,  $p$ 's  $> 0.19$ ).

The above contrasts suggest that the hippocampal head may selectively generalize across items that share both similar item and context information. However, it is possible that generalization within the hippocampus was graded, such that item-in-context generalization actually reflected the additive coding of item and context information separately. To arbitrate between additive (graded) and conjunctive (thresholded) accounts, pattern similarity estimates from hippocampal head were fit to a set of linear models, controlling for

hemisphere: an additive model (*both different* = 0, *similar item* = 1, *similar context* = 1, *similar item+context* = 2), a conjunctive model (*both different* = 0, *similar item* = 0, *similar context* = 0, *similar item+context* = 1), and a combined model containing both additive and conjunctive terms. Model fit was quantified using Bayesian Information Criterion (BIC; lower BIC values indicate better model fit) (Schwarz, 1978) and compared between the three models. BIC was slightly lower (indicating a better fit) for the conjunctive model (BIC = -952.36) than the additive model (BIC = -950.69). BIC was highest for the combined model (BIC = -942.63), suggesting that this model tended to overfit the data. These results suggest that the current pattern of hippocampal results is most consistent with conjunctive coding. Though additive coding cannot be conclusively ruled out on the basis of this analysis, there is no evidence for additive coding above what can be explained by conjunctive coding in our data.

Finally, we conducted a control analysis to determine whether pattern similarity in hippocampal head could be driven by the similarity of motor responses between trials, independent of memory for context information. Evoked patterns of activity across voxels in hippocampal head ROIs were correlated for each pair of trials given either a “Familiar” response (trials characterized by item familiarity without context recollection) or “New” response (forgotten trials). If pattern similarity was merely sensitive to motor response rather than episodic memory details, similarity estimates between trial pairs containing two “Familiar” or two “New” trials (i.e., *within* trials with the same motor response) would be expected to be higher than between trial pairs containing one “Familiar” and one “New” trial (i.e., *between* trials with different motor responses). However, a repeated measures ANOVA on response similarity, controlling for hemisphere, revealed no significant effects ( $F$ 's < 1.89,  $p$ 's > 0.18), suggesting that hippocampal pattern similarity was not driven by specific motor responses.

## DISCUSSION

In the current study, we used multivoxel pattern similarity analysis to investigate event representation in the hippocampus and MTL cortical areas during episodic retrieval. Anterior hippocampal voxel activity patterns were similar for events that involved similar items that had been associated with the same study context, but not for events with similar item or context information in isolation. These findings provide novel insight into the neural basis of memory for episodic details, suggesting that although neural coding in the hippocampus may differentiate between events with some overlapping attributes, it may generalize across events that share similar item-context relations.

Studies of BOLD signal magnitude (Bakker et al., 2008, Lacy et al., 2011) and multivariate voxel patterns (Liang et al., 2012, LaRocque et al., 2013, Huffman and Stark, 2014) have shown that, compared to MTL cortical regions, the hippocampus (particularly a combined ROI consisting of the dentate gyrus, CA2, and CA3) sharply differentiates between similar items. In the current study, we also found that the hippocampus represented similar items distinctly, as long as they had different contextual associations. However, when similar items shared contextual information, hippocampal activity patterns overlapped. This finding suggests an important and underexplored boundary condition on hippocampal pattern

separation: the extent to which the hippocampus differentiates between similar items may depend on the context in which the items had been encountered.

The present study used standard resolution fMRI, and it is possible that a richer pattern of results might have been revealed with high-resolution data. Most prior multivariate pattern similarity studies using high-resolution fMRI have not reported reliable subfield-level differences in item-level pattern separation (LaRocque et al., 2013, Huffman and Stark, 2014, Liang et al., 2012), but a recent study from our group found distinct representational profiles in CA1 versus a combined dentate gyrus/CA2/CA3 region during retrieval of item-context associations (Dimsdale-Zucker et al., 2018). In that study, participants viewed sequences of visual objects presented in virtual reality videos, and they were scanned while performing a recognition test on the studied objects, in the absence of any contextual cues. Results showed that pattern similarity was higher in CA1 across pairs of different objects that were studied in the same context (i.e., the same video) than for pairs of objects that had been studied in different contexts, whereas the dentate gyrus/CA2/CA3 ROI showed the opposite pattern (i.e., strongly differentiating between objects studied in the same context). In the present study, hippocampal activity patterns, aggregated across the subfields, were insensitive to context similarity if the items differed from one another. Based on the results of Dimsdale-Zucker et al. (2018), we can speculate that this null effect might have been the result of aggregating across CA1 (which might be expected to generalize across different items studied in the same context) and dentate gyrus/CA2/CA3 (which might be expected to differentiate between similar items studied in the same context). The present results, in turn, also suggest that both CA1 and dentate gyrus/CA2/CA3 might generalize across similar items that were studied in the same context. Future experiments can be designed to test this prediction at the subfield level.

The current results add importantly to previous work on hippocampal coding of item and context information. Previous studies have consistently suggested that hippocampal representations of items are context-dependent, but have not manipulated the similarity of item and context information. Consistent with our results, neuronal ensemble recordings in rats (McKenzie et al., 2014, Manns and Eichenbaum, 2009) have shown that the stability of hippocampal coding for specific items across repeated exposures is dependent on spatial context. Also in line with present findings, a previous fMRI study found that voxel patterns in the anterior hippocampus were more consistent across pairs of trials characterized by successful reinstatement of bound item-in-context information, compared to trials without associative retrieval (Hannula et al., 2013). The current study focused only on trials characterized by associative retrieval, demonstrating that item-in-context coding in the hippocampus is not merely a function of memory quality and is sensitive to the content of individual episodic memories. Additionally, our findings accord with recent fMRI studies in domains outside of episodic memory that relate the fidelity of hippocampal voxel patterns to implicit memory for the temporal position of items in learned sequences (Hsieh et al., 2014) and to accurate working memory for spatial configurations of items (Libby et al., 2014). Together, these results suggest that a key function of the hippocampus is tracking conjunctions of items and their temporal or spatial contexts, and that this phenomenon may override another key function – pattern separation – under certain conditions.

It is noteworthy that we observed similarity in hippocampal voxel patterns across *similar item+context* trial pairs, despite the fact that the corresponding study events were not exact duplicates. The events were similar at a schematic level but differed in their exact features. It could be argued that our results accord with the Complementary Learning Systems (CLS; Norman and O'Reilly, 2003, Norman, 2010) computational model, which predicts that hippocampal pattern separation breaks down when the average overlap across events is high. According to this view, however, we would expect participants to exhibit poor discriminability between related events (Elfman et al., 2008, Norman, 2010, Norman and O'Reilly, 2003), whereas item and context discriminability were very high in the current study. High memory performance cannot be explained in terms of schema-based retrieval (e.g., "I remember that there was a dog in the park"), because similar exemplars from each category were encoded in each context, and these items had to be discriminated from similar unstudied exemplars. Accordingly, participants had to use detailed representations to overcome interference between related events. Moreover, we did not see evidence of graded similarity of hippocampal representations, as might be expected based on reported simulations with the CLS model (Norman and O'Reilly, 2003, Norman, 2010). For instance, we saw no evidence for higher hippocampal pattern similarity for *similar item* than for *both different* pairs, and model comparisons suggested that hippocampal coding was better characterized as the conjunction of item and context information than the additive combination of individual features. The present results therefore suggest that models like CLS may benefit from explicitly incorporating representations of item and context information. If the model were instantiated such that contextual features play a significant and explicit role in hippocampal representations, then we would expect that the CLS model would exhibit pattern completion during processing of similar items encountered in the similar context, and pattern separation during processing of similar items encountered in different contexts.

Several recent studies have reported a role for the hippocampus in the integration of separate events that share key conceptual information (Chadwick et al., 2011, Horner et al., 2015, Schlichting et al., 2015, Milivojevic et al., 2015). The present results suggest that the similarity of item-context relations across events may be an important underlying factor that determines hippocampal integration. For instance, Milivojevic and colleagues (2015) showed that hippocampal voxel patterns are more similar across seemingly unrelated narrative videos when an intervening video reveals a link between the people (items) and locations (contexts) in the videos. On the basis of the current results, we would predict that an intervening video that shared context or items with the surrounding episodes, but not both, would disrupt hippocampal integration, perhaps particularly in anterior hippocampus. Indeed, in studies of complex pairwise item associations learned over the course of multiple trials, activity (Horner et al., 2015) and pattern similarity (Schlichting et al., 2015) in anterior hippocampus were greatest when event elements were fully integrated into "holistic" representations after learning. Given that the present study investigated single-shot episodic encoding and retrieval, our results suggest that anterior hippocampus integration or generalization can occur rapidly when item-context relations are sufficiently conceptually overlapping.

Everyday experiences often involve many common elements, yet we are able to retrieve specific instances of item-context associations – to which rack you locked your bike this afternoon, or which dog you saw in the park over the weekend – with surprising accuracy (Johnson et al., 1993, Mitchell and Johnson, 2009). Results from the current study suggest that the hippocampus utilizes context as an organizing principle in order to reduce competition between memories that involve the same or similar items. These findings bridge two influential theoretical frameworks on hippocampal function (Marr, 1971, Rolls and Kesner, 2006, Yassa and Stark, 2011, Norman and O’Reilly, 2003, Norman, 2010, Ranganath and Ritchey, 2012, Ritchey et al., 2015a, Eacott and Gaffan, 2005, Davachi, 2006, Ranganath, 2010, Knierim et al., 2006), by suggesting a critical role for the binding of item and context information in hippocampal pattern separation and episodic memory.

## ACKNOWLEDGEMENTS

This project was supported by a Guggenheim Fellowship and by a Vannevar Bush Faculty Fellowship (Office of Naval Research Grant N00014-15-1-0033) from the U.S. Department of Defense. Any opinions, findings, and conclusions or recommendations expressed in this material are those of the author(s) and do not necessarily reflect the views of the Office of Naval Research or the U.S. Department of Defense. In memory of Dr. Michael H. Buonocore.

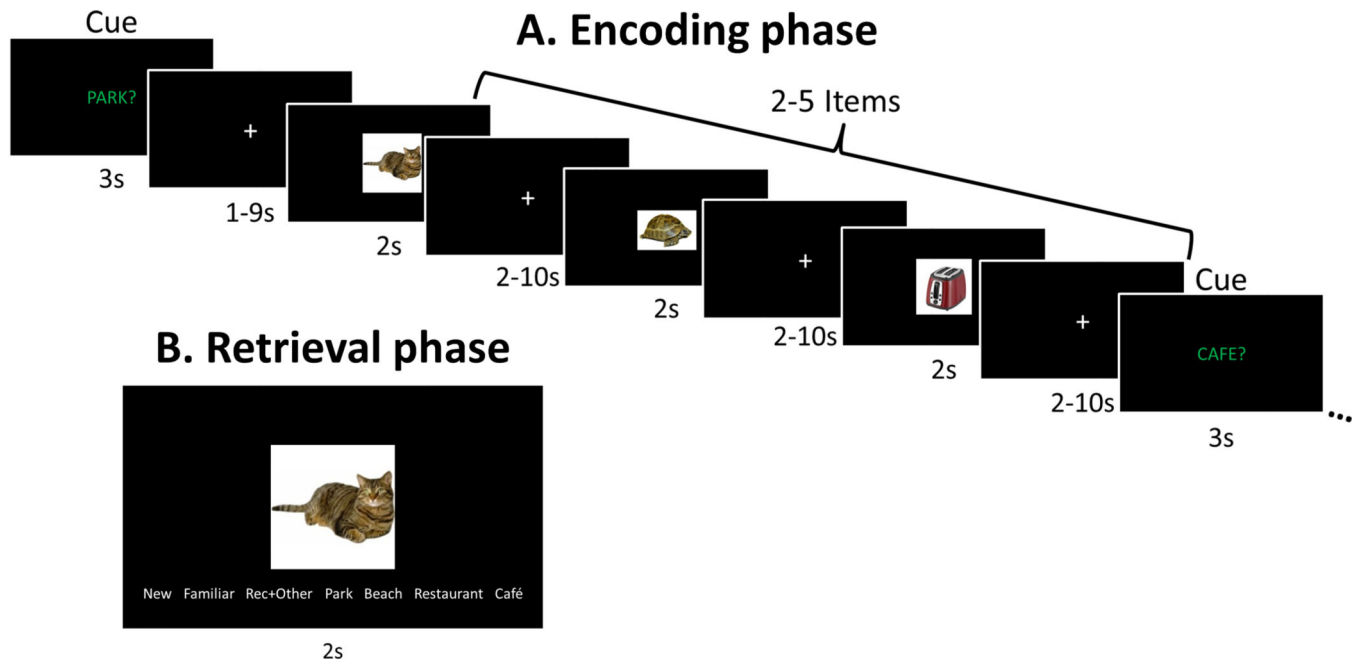
## REFERENCES

- BAKKER A, KIRWAN CB, MILLER M & STARK CE 2008 Pattern separation in the human hippocampal CA3 and dentate gyrus. *Science*, 319, 1640–2. [PubMed: 18356518]
- BERRON D, SCHÜTZE H, MAASS A, CARDENAS-BLANCO A, KUIJF HJ, KUMARAN D, & DÜZEL E 2016 Strong evidence for pattern separation in human dentate gyrus. *Journal of Neuroscience*, 36(29), 7569–7579. [PubMed: 27445136]
- BOSTOCK E, MULLER RU & KUBIE JL 1991 Experience-dependent modifications of hippocampal place cell firing. *Hippocampus*, 1, 193–205. [PubMed: 1669293]
- BROWN MW & AGGLETON JP 2001 Recognition memory: what are the roles of the perirhinal cortex and hippocampus? *Nat Rev Neurosci*, 2, 51–61. [PubMed: 11253359]
- BRUNEC IK, BELLANA B, OZUBKO JD, MAN V, ROBIN J, LIU ZX, GRADY C, ROSENBAUM RS, WINOCUR G, BARENSE M, & MOSCOVITCH M 2018 Multiple Scales of Representation along the Hippocampal Anteroposterior Axis in Humans. *Current Biology*, 28(13), 2129–2135. [PubMed: 29937352]
- CHADWICK MJ, HASSABIS D & MAGUIRE EA 2011 Decoding overlapping memories in the medial temporal lobes using high-resolution fMRI. *Learn Mem*, 18, 742–6. [PubMed: 22086391]
- COHEN NJ, POLDRACK RA & EICHENBAUM H 1997 Memory for items and memory for relations in the procedural/declarative memory framework. *Memory*, 5, 131–78. [PubMed: 9156097]
- DAVACHI L 2006 Item, context and relational episodic encoding in humans. *Curr Opin Neurobiol*, 16, 693–700. [PubMed: 17097284]
- DEUKER L, BELLMUND JL, SCHRÖDER TN, & DOELLER CF 2016 An event map of memory space in the hippocampus. *Elife*, 5, e16534. [PubMed: 27710766]
- DIMSDALE-ZUCKER HR, RITCHEY M, EKSTROM AD, YONELINAS AP, & RANGANATH C 2018 CA1 and CA3 differentially support spontaneous retrieval of episodic contexts within human hippocampal subfields. *Nature communications*, 9(1), 294.
- EACOTT MJ & GAFFAN EA 2005 The roles of perirhinal cortex, postrhinal cortex, and the fornix in memory for objects, contexts, and events in the rat. *Q J Exp Psychol B*, 58, 202–17. [PubMed: 16194965]
- ELFMAN KW, PARKS CM & YONELINAS AP 2008 Testing a neurocomputational model of recollection, familiarity, and source recognition. *J Exp Psychol Learn Mem Cogn*, 34, 752–68. [PubMed: 18605866]

- GUZOWSKI JF, KNIERIM JJ & MOSER EI 2004 Ensemble dynamics of hippocampal regions CA3 and CA1. *Neuron*, 44, 581–4. [PubMed: 15541306]
- HANNULA DE, LIBBY LA, YONELINAS AP & RANGANATH C 2013 Medial temporal lobe contributions to cued retrieval of items and contexts. *Neuropsychologia*.
- HORNER AJ, BISBY JA, BUSH D, LIN WJ & BURGESS N 2015 Evidence for holistic episodic recollection via hippocampal pattern completion. *Nat Commun*, 6, 7462. [PubMed: 26136141]
- HSIEH LT, GRUBER MJ, JENKINS LJ & RANGANATH C 2014 Hippocampal activity patterns carry information about objects in temporal context. *Neuron*, 81, 1165–78. [PubMed: 24607234]
- HUFFMAN DJ & STARK CE 2014 Multivariate pattern analysis of the human medial temporal lobe revealed representationally categorical cortex and representationally agnostic hippocampus. *Hippocampus*.
- JOHNSON MK, HASHTRUDI S & LINDSAY DS 1993 Source monitoring. *Psychol Bull*, 114, 3–28. [PubMed: 8346328]
- KNIERIM JJ, LEE I & HARGREAVES EL 2006 Hippocampal place cells: parallel input streams, subregional processing, and implications for episodic memory. *Hippocampus*, 16, 755–64. [PubMed: 16883558]
- KRIEGESKORTE N 2011 Pattern-information analysis: from stimulus decoding to computational-model testing. *Neuroimage*, 56, 411–21. [PubMed: 21281719]
- KRIEGESKORTE N, GOEBEL R & BANDETTINI P 2006 Information-based functional brain mapping. *Proc Natl Acad Sci U S A*, 103, 3863–8. [PubMed: 16537458]
- KRIEGESKORTE N, MUR M & BANDETTINI P 2008 Representational similarity analysis - connecting the branches of systems neuroscience. *Front Syst Neurosci*, 2, 4. [PubMed: 19104670]
- LACY JW, YASSA MA, STARK SM, MUFTULER LT & STARK CE 2011 Distinct pattern separation related transfer functions in human CA3/dentate and CA1 revealed using high-resolution fMRI and variable mnemonic similarity. *Learn Mem*, 18, 15–8. [PubMed: 21164173]
- LAROCQUE KF, SMITH ME, CARR VA, WITTHOFT N, GRILL-SPECTOR K & WAGNER AD 2013 Global similarity and pattern separation in the human medial temporal lobe predict subsequent memory. *J Neurosci*, 33, 5466–74. [PubMed: 23536062]
- LEUTGEB JK, LEUTGEB S, MOSER MB & MOSER EI 2007 Pattern separation in the dentate gyrus and CA3 of the hippocampus. *Science*, 315, 961–6. [PubMed: 17303747]
- LEVER C, WILLS T, CACUCCI F, BURGESS N & O'KEEFE J 2002 Long-term plasticity in hippocampal place-cell representation of environmental geometry. *Nature*, 416, 90–4. [PubMed: 11882899]
- LIANG JC, WAGNER AD & PRESTON AR 2012 Content Representation in the Human Medial Temporal Lobe. *Cereb Cortex*.
- LIBBY LA, HANNULA DE & RANGANATH C 2014 Medial Temporal Lobe Coding of Item and Spatial Information during Relational Binding in Working Memory. *J Neurosci*, 34, 14233–42. [PubMed: 25339737]
- MALDJIAN JA, LAURIENTI PJ, KRAFT RA & BURDETTE JH 2003 An automated method for neuroanatomic and cytoarchitectonic atlas-based interrogation of fMRI data sets. *Neuroimage*, 19, 1233–9. [PubMed: 12880848]
- MANN JR & EICHENBAUM H 2009 A cognitive map for object memory in the hippocampus. *Learn Mem*, 16, 616–24. [PubMed: 19794187]
- MARR D 1971 Simple memory: a theory for archicortex. *Philos Trans R Soc Lond B Biol Sci*, 262, 23–81. [PubMed: 4399412]
- MCKENZIE S, FRANK AJ, KINSKY NR, PORTER B, RIVIÈRE PD & EICHENBAUM H 2014 Hippocampal representation of related and opposing memories develop within distinct, hierarchically organized neural schemas. *Neuron*, 83, 202–15. [PubMed: 24910078]
- MILIVOJEVIC B, VICENTE-GRABOVETSKY A & DOELLER CF 2015 Insight reconfigures hippocampal-prefrontal memories. *Curr Biol*, 25, 821–30. [PubMed: 25728693]
- MITCHELL KJ & JOHNSON MK 2009 Source monitoring 15 years later: what have we learned from fMRI about the neural mechanisms of source memory? *Psychol Bull*, 135, 638–77. [PubMed: 19586165]

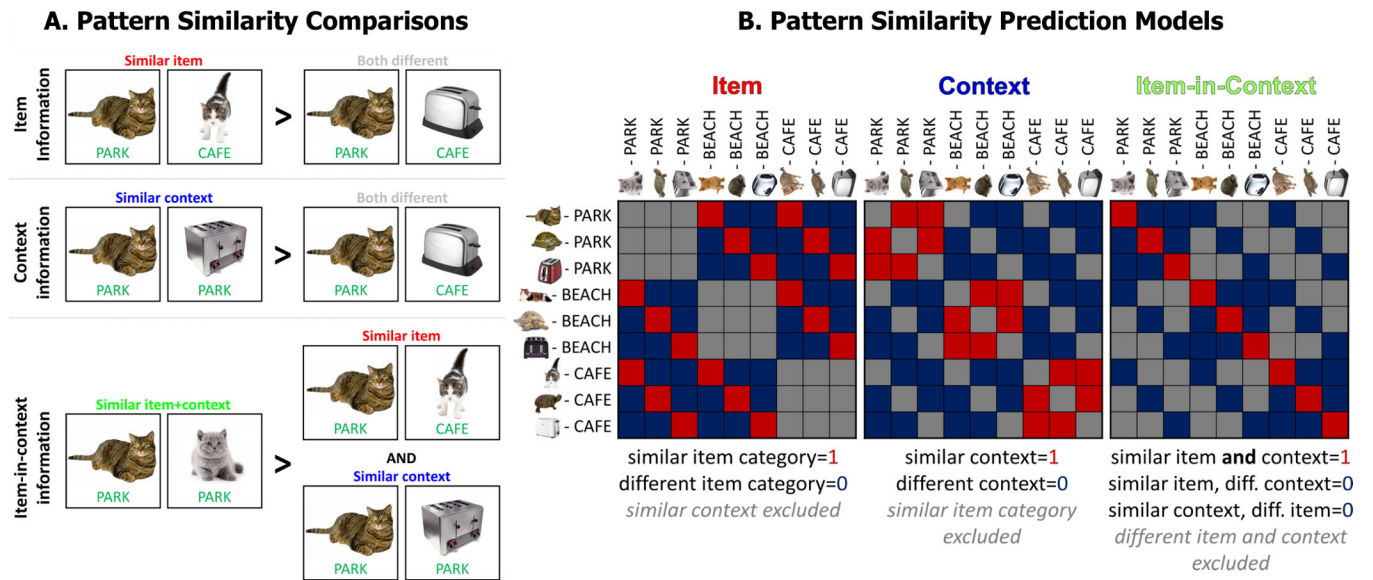


- MOORE M, HU Y, WOO S, O'HEARN D, IORDAN AD, DOLCOS S & DOLCOS F 2014 A comprehensive protocol for manual segmentation of the medial temporal lobe structures. *J Vis Exp*.
- MUMFORD JA, DAVIS T & POLDRACK RA 2014 The impact of study design on pattern estimation for single-trial multivariate pattern analysis. *Neuroimage*.
- MUMFORD JA, TURNER BO, ASHBY FG & POLDRACK RA 2012 Deconvolving BOLD activation in event-related designs for multivoxel pattern classification analyses. *Neuroimage*, 59, 2636–43. [PubMed: 21924359]
- NEUNUEBEL JP & KNIERIM JJ 2014 CA3 retrieves coherent representations from degraded input: direct evidence for CA3 pattern completion and dentate gyrus pattern separation. *Neuron*, 81(2), 416–427. [PubMed: 24462102]
- NICHOLS TE & HOLMES AP 2002 Nonparametric permutation tests for functional neuroimaging: a primer with examples. *Hum Brain Mapp*, 15, 1–25. [PubMed: 11747097]
- NORMAN KA 2010 How hippocampus and cortex contribute to recognition memory: revisiting the complementary learning systems model. *Hippocampus*, 20, 1217–27. [PubMed: 20857486]
- NORMAN KA & O'REILLY RC 2003 Modeling hippocampal and neocortical contributions to recognition memory: a complementary-learning-systems approach. *Psychol Rev*, 110, 611–46. [PubMed: 14599236]
- POPPEK J, EVENSMOEN HR, MOSCOVITCH M & NADEL L 2013 Long-axis specialization of the human hippocampus. *Trends Cogn Sci*, 17, 230–40. [PubMed: 23597720]
- RANGANATH C 2010 A unified framework for the functional organization of the medial temporal lobes and the phenomenology of episodic memory. *Hippocampus*, 20, 1263–90. [PubMed: 20928833]
- RANGANATH C & RITCHEY M 2012 Two cortical systems for memory-guided behaviour. *Nat Rev Neurosci*, 13, 713–26. [PubMed: 22992647]
- RITCHEY M, LIBBY LA & RANGANATH C 2015a Cortico-hippocampal systems involved in memory and cognition: the PMAT framework. *Prog Brain Res*, 219, 45–64. [PubMed: 26072233]
- RITCHEY M, MONTCHAL ME, YONELINAS AP & RANGANATH C 2015b Delay-dependent contributions of medial temporal lobe regions to episodic memory retrieval. *Elife*, 4.
- ROLLS ET & KESNER RP 2006 A computational theory of hippocampal function, and empirical tests of the theory. *Prog Neurobiol*, 79, 1–48. [PubMed: 16781044]
- SCHLICHTING ML, MUMFORD JA & PRESTON AR 2015 Learning-related representational changes reveal dissociable integration and separation signatures in the hippocampus and prefrontal cortex. *Nat Commun*, 6, 8151. [PubMed: 26303198]
- SCOVILLE WB & MILNER B 1957 Loss of recent memory after bilateral hippocampal lesions. *J Neurol Neurosurg Psychiatry*, 20, 11–21. [PubMed: 13406589]
- SHELDON S & LEVINE B 2015 The medial temporal lobes distinguish between within-item and item-context relations during autobiographical memory retrieval. *Hippocampus*, 25(12), 1577–1590. [PubMed: 26032447]
- VARGHA-KHADEM F, GADIAN DG, WATKINS KE, CONNELLY A, VAN PAESSCHEN W & MISHKIN M 1997 Differential effects of early hippocampal pathology on episodic and semantic memory. *Science*, 277, 376–80. [PubMed: 9219696]
- WILLS TJ, LEVER C, CACUCCI F, BURGESS N & O'KEEFE J 2005 Attractor dynamics in the hippocampal representation of the local environment. *Science*, 308, 873–6. [PubMed: 15879220]
- XUE G, DONG Q, CHEN C, LU Z, MUMFORD JA & POLDRACK RA 2010 Greater neural pattern similarity across repetitions is associated with better memory. *Science*, 330, 97–101. [PubMed: 20829453]
- YASSA MA & STARK CE 2011 Pattern separation in the hippocampus. *Trends Neurosci*, 34, 515–25. [PubMed: 21788086]

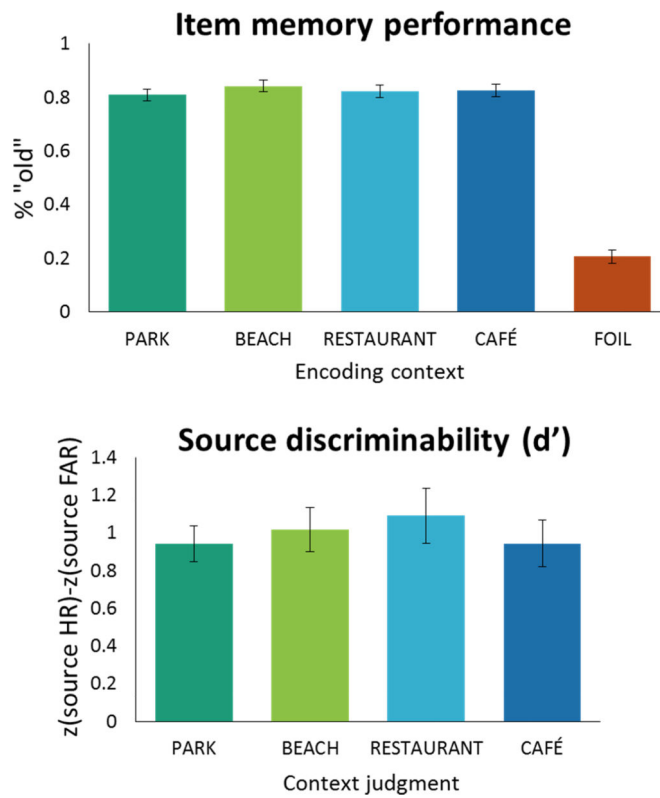


**Figure 1.**

Task design. A) During encoding, trial-unique exemplars from a set of categories were presented sequentially, broken down into mini-blocks separated by a cue screen indicating the encoding context. Two exemplars from each category were encoded in each context. B) At retrieval, all targets and a fully category-matched set of foils were presented in a combined recognition/cued recall test. Only trials given correct source memory judgments were included in voxel pattern similarity analyses.

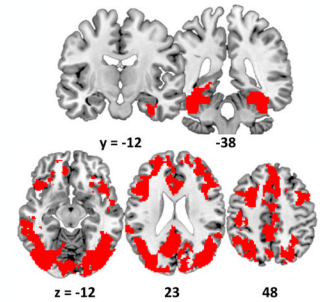
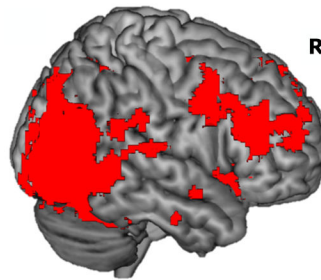
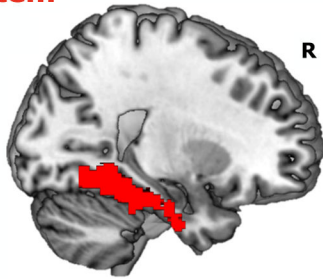
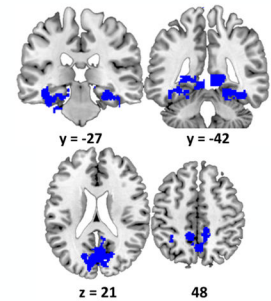
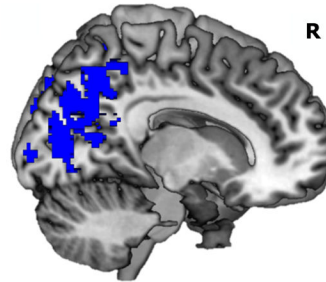
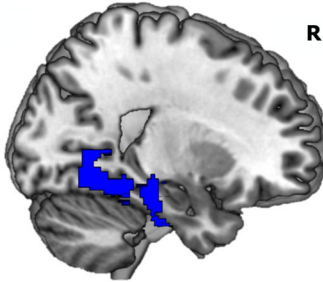
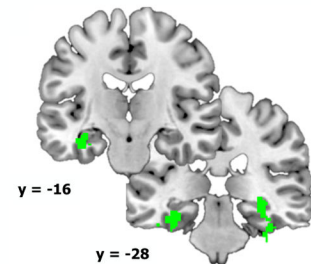
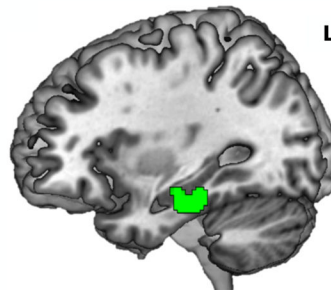
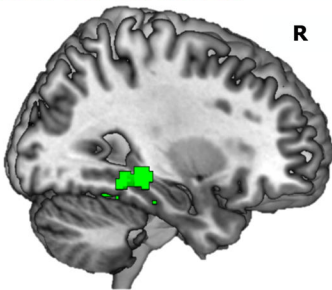


**Figure 2.** Trial pair similarity comparisons and models. A) Example trial pair organization for pattern similarity analyses. Item information coding was estimated by contrasting pattern similarity estimates between “similar item” and “both different” trial pairs. Context coding was estimated by contrasting “similar context” and “both different” trial pairs. Item-in-context coding was calculated by contrasting “similar item+context” trial pairs against both “similar item” and “similar context” pairs. B) Example trial-by-trial similarity model matrices. Item and context predicted similarity models were orthogonal to each other; the item-in-context predicted similarity model was hierarchical to the item and context models. Individual subject pattern similarity matrices were correlated (biserial) with model matrices to test for item, context, and item-in-context generalization.

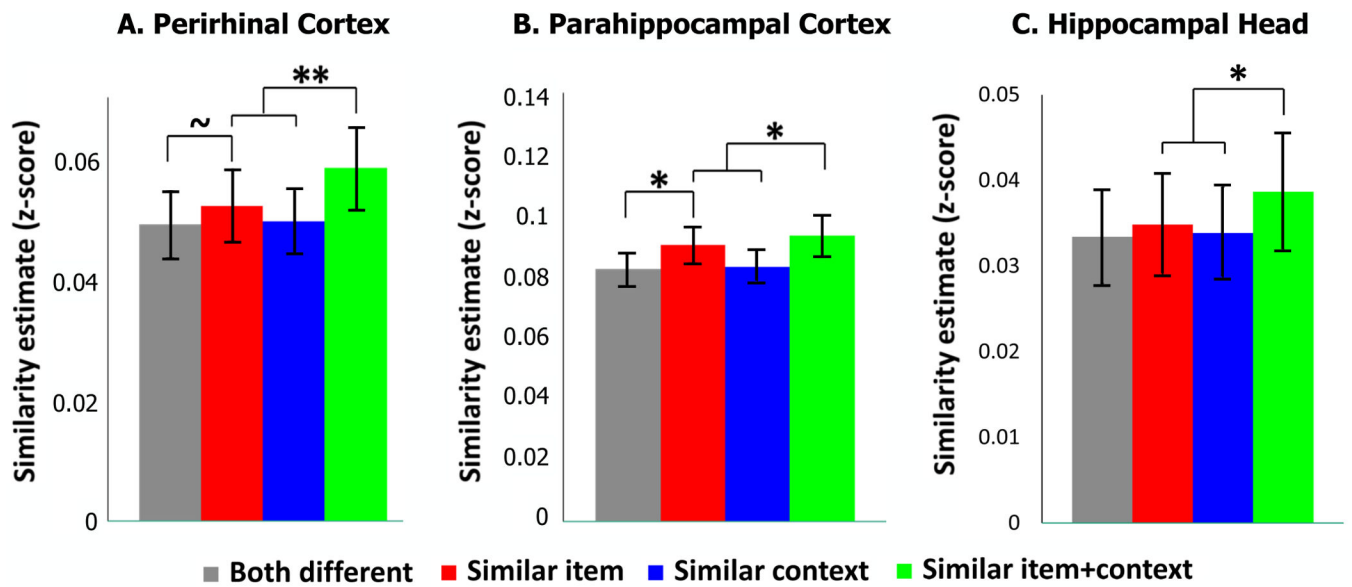


**Figure 3.**

Behavioral expressions of item and context memory were equivalent across encoding context locations. Item memory performance (top) was calculated as the proportion of old items from each encoding context given any “old” response (“Familiar,” “Recollect,” or any source judgment); item false alarm rate was calculated as the proportion of foils given any “old” response. Source discriminability (bottom) was calculated as the difference between the inverse normal cumulative distributions of source hit rate (HR) and source false alarm rate (FAR). For each context judgment, source HR was calculated as the proportion of items correctly attributed to a particular encoding condition; source FAR was calculated as the proportion of items (targets or foils) incorrectly attributed to that encoding condition. Error bars indicate standard error of the mean across participants.

**A. Item****B. Context****C. Item-In-Context****Figure 4.**

Voxel pattern similarity analysis results. A) *similar item > both different* contrast. Using an MTL mask, significant clusters were identified in ventral and medial temporal cortex, including perirhinal cortex. Using a whole-brain grey matter mask (excluding MTL voxels; middle), extensive significant clusters were identified, with the global maximum in posterior fusiform cortex (MNI 40, -56, -14;  $t = 10.22$ ). B) Association between *similar context > both different* pattern similarity and context memory performance. With an MTL mask, the global maximum in parahippocampal cortex (MNI 28, -24, -20;  $t = 6.61$ ) was significant at  $p_{TFCE} < 0.05$ , FWE-corrected (for display purposes, threshold is set to  $p < 0.001$ , uncorrected). With a whole-brain grey matter mask (excluding MTL voxels; middle), significant clusters were identified in posterior medial regions, with the global maximum in cuneus (MNI -10, -86, 16;  $t = 6.82$ ). C) Searchlight results for *similar item+context > (similar item or similar context)* contrast in sagittal and coronal views. Thresholds set to  $p < 0.05$ , FWE-corrected.

**Figure 5.**

Pattern similarity results across MTL ROIs. Mean pattern similarity estimates for both different, similar item, similar context, and similar item+context trial pairs are plotted. A) In Perirhinal Cortex, there was evidence for both item and item-in-context generalization. B) In Parahippocampal Cortex, item-in-context generalization was redundant with item coding effects. C) In the hippocampal head, there was evidence for generalization due to item-in-context associations, but not item or context similarity individually.  $\sim p \approx 0.05$  (trend-level), \*  $p < 0.05$  for a priori contrasts (indicated by brackets). \*\*  $p < 0.05$  for both a priori and post-hoc (similar item+context > similar item) contrasts. Error bars refer to standard error across participants. Because there was no significant differences across hemisphere, results are collapsed across right and left ROIs.