

The neural correlates of gist-based true and false recognition

Angela H. Gutchess^{a,b,*}, Daniel L. Schacter^c

^a Brandeis University, USA

^b Massachusetts General Hospital, USA

^c Harvard University, USA

ARTICLE INFO

Article history:

Received 21 July 2011

Revised 25 November 2011

Accepted 26 November 2011

Available online 8 December 2011

Keywords:

Memory

False recognition

Gist

fMRI

Hippocampus

ABSTRACT

When information is thematically related to previously studied information, gist-based processes contribute to false recognition. Using functional MRI, we examined the neural correlates of gist-based recognition as a function of increasing numbers of studied exemplars. Sixteen participants incidentally encoded small, medium, and large sets of pictures, and we compared the neural response at recognition using parametric modulation analyses. For hits, regions in middle occipital, middle temporal, and posterior parietal cortex linearly modulated their activity according to the number of related encoded items. For false alarms, visual, parietal, and hippocampal regions were modulated as a function of the encoded set size. The present results are consistent with prior work in that the neural regions supporting veridical memory also contribute to false memory for related information. The results also reveal that these regions respond to the degree of relatedness among similar items, and implicate perceptual and constructive processes in gist-based false memory.

© 2011 Elsevier Inc. All rights reserved.

Introduction

Memory is a constructive process that is sometimes prone to error and distortion (e.g., Bartlett, 1932; Loftus, 1979; Neisser, 1967; Schacter, 2001; Schacter and Addis, 2007; Schacter et al., 1998). The notion that memory distortions are a byproduct of a constructive memory system is supported by a wealth of behavioral data revealing the nature of these errors (e.g., Brainerd and Reyna, 2002; Deese, 1959; Mather et al., 1997; Roediger and McDermott, 1995, 2000; Schacter et al., 1999) (for reviews see Gallo, 2006, 2010; Newman and Lindsay, 2009; Schacter et al., 2011). For example, in his classic studies, Bartlett (1932) provided evidence that errors in story recall were frequently attributable to the operation of schemas that help to organize experiences and guide retrieval processes. Other memory errors reflect the operation of associative memory processes that play a key role in organizing memory or related gist-based processes that promote the retention of thematic information (Deese, 1959; Reyna and Brainerd, 1995; Roediger and McDermott, 1995). These kinds of findings have provided empirical support for the claim that a range of memory distortions is based on the operation of adaptive cognitive processes that contribute to the efficient and flexible functioning of memory (for further discussion, see Newman and Lindsay, 2009; Schacter and Addis, 2007; Schacter et al., 2011).

Neuroimaging studies of *false recognition*, where individuals incorrectly claim that a novel item has been encountered previously, reveal

a striking overlap between the brain regions activated during true and false recognition (e.g., Abe et al., 2008; Cabeza et al., 2001; Dennis et al., 2008; Garoff-Eaton et al., 2006; Johnson et al., 1997; Kensinger and Schacter, 2006; Okado and Stark, 2003; Schacter et al., 1997; Schacter et al., 1996; Slotnick and Schacter, 2004); (see review by Schacter et al., *in press*; Schacter and Slotnick, 2004). Such observations are consistent with the idea that when memory errors occur, there is a high degree of overlap in the subprocesses that contribute to true and false memory (Mitchell and Johnson, 2009), and provide additional evidence supporting claims of adaptive memory distortions (Schacter et al., 2011).

Previous fMRI studies examining false recognition of novel items that are related to previously studied items indicates the types of overlapping processes that can lead to false recognition. In a comparison of true recognition, related false recognition (e.g., of perceptually similar novel shapes), and unrelated false recognition (e.g., perceptually dissimilar novel shapes), Garoff-Eaton et al. (2006) noted that a wide array of regions, including prefrontal, lateral and medial temporal, parietal, and occipital cortices, was activated by both identical true and related false recognition. These findings suggest that related false recognition engages many of the same processes as accurate recognition, including semantic and visual processes. Studies of associative memory errors also suggest that semantic processes contribute to false memory, with heightened engagement of inferior frontal gyrus during encoding (Kim and Cabeza, 2007) and retrieval (Garoff-Eaton et al., 2007), and a possible role for anterior temporal regions that subserve semantic processing (Gallate et al., 2009). Perceptual regions implicated in false memory include late visual regions, which respond to both true and false memory (Kim and

* Corresponding author at: Brandeis University, Department of Psychology, MS 062, P.O. Box 549110, Waltham, MA 02454-9110, USA. Fax: +1 781 736 3291.

E-mail address: gutchess@brandeis.edu (A.H. Gutchess).

Cabeza, 2007; Slotnick and Schacter, 2004), and a midtemporal region responding more for false than true memories for rhyming or orthographically similar words (Garoff-Eaton, et al., 2007).

The paradigms employed thus far, however, have not investigated false recognition based on the *degree* or *amount* of related information that is available at the time of retrieval (see Kim and Cabeza, 2007, and Dennis et al., 2007, for studies of encoding activity based on the extent of false recognition of categorized words). While several neural regions are engaged when there is some degree of overlap between novel and familiar exemplars, there may only be a subset of regions that are increasingly engaged as the gist-based representation is strengthened. Characterizing and understanding these regions is particularly relevant to the claim that memory distortions often reflect the operation of adaptive cognitive processes (Schacter et al., 2011). In this study, we explore the neural regions that respond during true and false recognition as a function of the number of related items. Furthermore, the present paradigm incorporates visually and conceptually rich pictures, which should draw on both perceptual and semantic processes, in contrast to previous studies (see Dennis, et al., 2008; Dennis et al., 2007; Kim and Cabeza, 2007 for comparisons of categorically-related words).

To investigate the effects of the degree of relatedness on neural activity during false recognition, we focus on a memory error known as *gist-based false recognition*. Gist-based errors occurs when people extract the gist, or general information about thematic content, but fail to encode or retrieve verbatim, item-specific distinguishing details (Brainerd and Reyna, 2002; Koutstaal and Schacter, 1997). Strengthening gist representations by encountering multiple exemplars of a class of objects can lead to false recognition of items thematically related to studied exemplars (Koutstaal and Schacter, 1997). Importantly, the number of memory errors seems to be driven by the amount of related information; as the number of related studied items increases, so too does the percentage of items falsely recognized (Koutstaal and Schacter, 1997). For example, studying a dozen pictures of cats leads one to remember quite well that cats were studied, but can hinder one's ability to recall perceptual details of particular cats and thus to discriminate novel pictures of cats from studied pictures of cats. Stimulus characteristics, instructions, and test conditions all can influence reliance on gist-based responding (Guerin et al., in press; Koutstaal and Cavendish, 2006; Koutstaal et al., 1999; Koutstaal et al., 2003; Tun et al., 1998).

A number of cognitive processes have been proposed to contribute to gist-based memory errors. The semantic categorization account maintains that pre-existing semantic associations can overshadow the use of item-specific features, and is predicated on the finding that studying sets of concrete, but not abstract, pictures increases levels of false recognition (Koutstaal, et al., 2003). Retrieval processes contribute substantially to gist-based errors (Guerin et al., in press), as disambiguating perceptual information is adequately encoded to at least some extent (Koutstaal, 2003), and can be made accessible under appropriate retrieval conditions (Guerin et al., in press). In addition, encountering many overlapping perceptual features can reinforce the representation of prototypical features (e.g., the shape features that are generally shared across exemplars from the category "chair") which then, as a result of flexible recombination or binding of features by a constructive memory system, could seem familiar in a novel exemplar that shares those same prototypical features.

In addition to semantic and perceptual processes, retrieval also relies on reconstructive processes to combine information from various sources, and the hippocampus appears to contribute in this role. For example, the hippocampus is engaged more by the correct recognition of a previously encountered word than by a novel recombination of parts of the word in young adults, and is implicated in age-related failures to correctly bind features together (Giovanello et al., 2010; Mitchell et al., 2000). Recent work suggests that the region's role in connecting information together also contributes to imagination.

The hippocampus is engaged both by imagining a future scenario and retrieving a memory of the past (e.g., Addis et al., 2007). Constructing future scenarios can even drive the region more than retrieving memories of the past (Schacter and Addis, 2007), particularly engaging anterior hippocampal regions when constructed events contain high levels of detail (Addis and Schacter, 2008) and reflect specific, rather than general, scenarios (Addis et al., 2011). This role for the hippocampus is thought to reflect the relational processing demands of integrating and recombining information from a variety of sources to imagine and encode a new scenario (Addis et al., 2007, 2009; Hassabis et al., 2007; Martin et al., 2011) (for review and discussion, see Buckner, 2010; Schacter and Addis, 2009).

Consistent with prior studies of false recognition of related information, we expect that gist-based errors will occur due to the engagement of semantic, perceptual, and constructive processes. However, investigating false recognition as a function of gist strength, based on having encountered previously varying numbers of related items, allows for the investigation of false memory across gradations of relatedness. While previous studies have begun to identify shared neural substrates for accurate memory and erroneous memory for related information, relatedness has been treated in an all-or-none fashion. It may be that a minimal level of relatedness is all that is required to invoke similar processes to support true and false recognition, in which case we should find that perceptual, semantic, and constructive processes are no more engaged as gist strength increases. However, behavioral data indicating higher levels of gist-based false recognition as a function of the number of related items studied (Koutstaal and Schacter, 1997) suggest that these processes and their corresponding neural regions should be increasingly engaged as function of gist strength.

Our approach based on manipulating gist strength should be particularly useful for shedding light on the mixed results in the literature thus far, with some studies indicating that the hippocampus and other regions are engaged equally by true and false memories (e.g., Addis, et al., 2007; Garoff-Eaton, et al., 2006; Schacter, et al., 1997), while some suggest that the regions are engaged more by true than false memories (e.g., Cabeza, et al., 2001; Giovanello, et al., 2010; Kensinger and Schacter, 2005; Schacter, et al., 1996), or even more by imaginary than true memories, as in the research on future event simulation (e.g., Addis, et al., 2007; Schacter and Addis, 2007). One possibility is that the hippocampus responds to the *degree* of relatedness among similar items, such that false memories that have more information in common with true ones implicate these regions more than those false memories with less in common with true memories. We tested the response of the hippocampus and other regions by manipulating the strength of the gist-based representation for conceptually and perceptually rich information. Using a parametric modulation approach, we identify the neural regions that respond as the number of related items increases, allowing us to selectively identify those cognitive processes leading to increased levels of gist-based recognition.

Methods

Participants

Sixteen participants (8 male) between the ages of 19–33 ($M=24.13$, $SD=4.57$) were recruited to take part in the study. Seven additional participants were excluded from the study for failing to respond to large numbers of trials (1 participant; >40% no responses), failure to follow instructions (1 participant), or too few false alarms in at least one condition (5 participants; <8 items). The final sample of participants averaged 16.41 years of education ($SD=2.85$), and all participants had some college education. Eligibility criteria included right handedness, English as a native language, absence from medications known to affect the central nervous

system, and the absence of neurological, psychological, or physical conditions contraindicated for MRI scanning. Participants provided written informed consent for a protocol approved by Harvard University and Partners Institutional Review Board.

Materials and procedures

Participants incidentally encoded 468 pictures of single objects by making a yes/no button press to denote whether the type of object depicted is something they would use or interact with during an average day. Pictures were drawn from 54 sets of objects (e.g., umbrellas, chairs, cats), with 18 categories assigned to each condition. Multiple exemplars of each object were encoded with small study sets containing four studied exemplars (e.g., four umbrellas), medium study sets containing eight studied exemplars (e.g., eight chairs), and large studied sets containing fourteen exemplars (e.g., fourteen cats). Exemplars from each category were distributed across three encoding runs (e.g., the four umbrellas or fourteen cats would be distributed as evenly as possible across encoding runs). Across three encoding runs, participants viewed 72 pictures that would ultimately comprise the small condition, 144 medium set pictures, and 252 large set pictures. Each picture was presented for one second followed by a 1-second blank interval. On average, participants responded to 94.24% of the trials during encoding ($SD = 8.20\%$).

After an approximately ten minute delay during which time structural images were acquired, participants received a surprise recognition test. Participants' memory was assessed for a total of 456 pictures (216 studied items and 240 lures). Each studied small, medium, and large object category (e.g., cats) was tested with four studied exemplars and four novel lure exemplars, randomly selected from the studied and unstudied items for each category. Thus, there was a total of 72 items in each of the conditions, with items from each specific category from each condition evenly distributed across four runs. In addition, twenty-four novel pictures were presented from distinct object classes not studied previously in order to provide a baseline false alarm rate. Example stimuli are displayed in Fig. 1. Counterbalancing of the assignment of object classes to each of the

four studied set sizes (i.e., zero, small, medium, and large) and assignment of tested items as either lures or studied targets was accomplished across participants using eight different recognition orders. Participants had four seconds in which to decide whether or not they had previously seen the picture and to press a button corresponding to “yes” or “no”. Pictures were selected from photo CDs (Hemera Technologies, Gatineau, Quebec).

Both encoding and recognition trials were randomly ordered in a jittered design (Dale, 1999) containing fixation cross trials that participants passively viewed for times varying from 2000 to 10000 ms. Trials were presented and behavioral data acquired with E-Prime software (Psychology Software Tools, Pittsburgh, PA).

Functional MRI data acquisition

Images were acquired using a Siemens Avanto 1.5 Tesla whole-body scanner (Siemens, Erlangen, Germany). Thirty-two slices 3.2 mm thick with a .3 mm skip between slices were acquired with an echo-planar imaging (EPI) sequence ($TR = 2000$ ms, $TE = 30$, $FOV = 200$ mm, and a flip angle = 90°). During each of the three encoding runs, 212 measurements were collected; each of the four recognition runs consisted of 304 measurements. Encoding data are intended for a separate investigation and will not be discussed further.

Functional MRI analyses

Analyses were conducted in SPM2 (Wellcome Department of Cognitive Neurology, London, UK). Random effects analyses allowed for assessment of distinct and common patterns of activation, and parametric analyses were conducted to identify regions that modulate neural activity during hits or false alarms as an effect of the amount of gist. The random effects model included eight regressors: False alarms (FA)—large, FA—medium, FA—small, Hits—large, Hits—medium, Hits—small, Zero—Correct Rejections (novel lures), and Miscellaneous (i.e., all misses and remaining correct rejections; when applicable, false alarms to the Zero category and non-



Fig. 1. Example stimuli are displayed for a medium set size. Participants encoded eight exemplars (outlined in yellow). At the time of recognition, they were tested on four of the previously presented exemplars (middle row, outlined in yellow and green) as well as four new lure exemplars (bottom row, outlined in green). Four old and four new exemplars were tested at recognition for all set sizes, although the number of encoded exemplars ranged from four (i.e., small sets) to fourteen (i.e., large sets).

response trials). Results are thresholded at a voxel-level correction of $p < .01$ with a cluster-level threshold of 44 voxels in order to achieve an overall correction of $p < .05$ (Slotnick et al., 2003). To assess the commonality, or overlap, across two comparisons, two separate contrasts were estimated. The map for the first contrast was thresholded at $p < .035$ (Fisher, 1950; Garoff-Eaton, et al., 2006) to create a mask in which the second contrast was tested at a threshold of $p < .035$ (for a conjoint p-value of approximately $p = .001$) and a cluster extent = 44 voxels (for consistency with the difference analyses), for an overall correction surpassing $p < .05$.

The parametric modulation analyses were conducted at the fixed-effects level, entering set size (i.e., small, medium, and large) as a covariate of interest, to test for linear trends across set sizes separately for hits and false alarms. These analyses allowed for the identification of regions whose activity correlated positively (i.e., increasing as set size increased) and negatively (i.e., increasing as set size decreased) modulations across the different levels of set size for either hits or false alarms. Because the parametric modulation responses could differ depending on whether recognition was true or false, we directly compared the slope of the parametric responses across these two conditions by entering the positive contrast images from the fixed-effects models into random-effects paired samples t-tests in SPM (i.e., testing whether the positive modulation effect differed between true and false recognition, and whether the negative modulation effect differed between true and false recognition) (as in Addis and Schacter, 2008). Any slope differences identified between conditions could reflect: (1) slopes of the same sign that differ significantly in magnitude across conditions (e.g., the parametric effect is strongly positive for true recognition and weakly positive for false recognition); or (2) slopes of opposite sign (e.g., the parametric effect is positive for true recognition and negative for false recognition). Moreover, it is possible that even when there are significant differences in slope across conditions in a particular voxel, the parametric effects themselves may not be significantly different from zero (e.g., a non-significant positive effect for true recognition and a weak non-significant negative for false recognition). Thus, in order to fully explore all these possibilities, we computed a random-effects one-sample t-test for the parametric effect in each condition. The values in the resultant beta images from these one-sample t-tests reflect the average estimated slope of the regression line in each condition; these beta images could then be probed (in regions identified by the paired sample t-test as showing a differing parametric effect across condition) to examine the nature of the parametric effect (sign; significance; see Table 3).

For the purposes of illustration, activations are displayed on a canonical single subject's brain. Estimates for bar graphs are based on random effects analyses with values extracted with MarsBar (Brett et al., 2002). Labels of regions are approximate, based on Talairach (Talairach and Tournoux, 1988) and AAL labels (Tzourio-Mazoyer et al., 2002), as implemented in mricron (Rorden and Brett, 2000).

Results

Behavioral recognition data

Participants showed the expected pattern of greater false alarms as a function of increasing set size, as tested in a univariate ANOVA of false alarm rates across the three previously studied set sizes $F(2, 30) = 17.67$, $p < .001$, partial $\eta^2 = .54$. Each set size differed significantly from the others, with more false alarms to large than medium set sizes, $F(1, 15) = 11.07$, $p < .01$, partial $\eta^2 = .43$, and to medium than small, $F(1, 15) = 8.90$, $p < .01$, partial $\eta^2 = .37$. Participants made fewer false alarms to the entirely novel zero items than to any other set, $t(30) = 11.28$, $p < .001$ (for the comparison against the small set). In contrast to the false alarm rate data, set size did not influence hit rates across the three different set sizes when tested in a univariate ANOVA, $F(1, 15) = 1.78$, $p = .19$, partial $\eta^2 = .11$ See Table 1.

Table 1

This table summarizes the behavioral data (means and standard deviations) for hit and false alarm rates for each set size.

	Zero	Small	Medium	Large
Hits	N/A	.60 (.14)	.63 (.15)	.63 (.12)
False alarms	.04 (.07)	.19 (.09)	.24 (.13)	.30 (.10)

Functional data: random effects analyses

Common activity for true and false recognition

We first assessed the overlap of the activations for hits and false alarms as a baseline comparison of whether the overlap reported in previous studies for true and false recognition was also true for our study. The conjunction of the hits minus novel items (i.e., encoded set size of zero) contrast and the false alarms minus novel items contrast resulted in several regions of activation, including bilateral

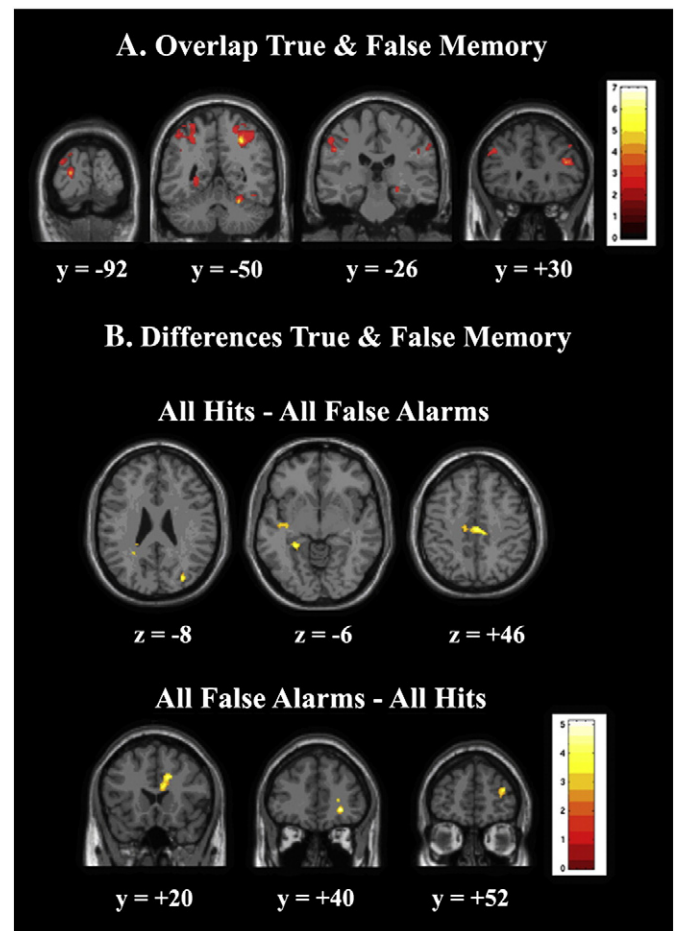


Fig. 2. A. Overlap between true and false recognition (compared to novel items) occurs in a variety of regions. The selected slices (from left to right) display activations in left middle occipital (peak = $-20, -92, 10$), bilateral inferior parietal (right peak = $28, -50, 48$; left peak = $-28, -46, 48$), right hippocampal ($26, -26, -10$), and bilateral inferior frontal (right peak = $42, 30, 20$; left peak = $-42, 24, 30$) regions. B. Differences between true and false recognition (collapsed across set size) occur in predominantly sensory regions for the comparison of hits–false alarms and in predominantly frontal control regions for the comparison of false alarms–hits (bottom row). Activations for the comparison of true minus false recognition (displayed on the top row) include right middle occipital (peak = $32, -80, 26$), left parahippocampal ($-26, -40, -6$), left middle temporal ($-50, -10, -8$) and left posterior parietal (peak = $4, -20, 46$) cortex. Activations for the false alarms minus hits comparison (displayed on the bottom row) include right anterior cingulate (peak = $16, 20, 34$), two regions of right middle frontal gyrus (ventral peak = $30, 40, -4$; dorsal peak = $36, 52, 18$).

inferior parietal, bilateral inferior and middle frontal, bilateral precuneus, and occipital gyri, as well as right hippocampus. A subset of these regions is displayed in Fig. 2A (the full set of coordinates is available as Supplementary Table 1).

Differences in the activity for true and false recognition

In a comparison collapsing across set sizes, we tested for differences in the activity underlying true and false recognition. Representative slices from these comparisons are displayed in Fig. 2B (the full set of coordinates is available as Supplementary Table 2). Hits tended to recruit visual, parahippocampal, and temporal regions more than false alarms, whereas false alarms recruited regions of frontal cortex, including anterior cingulate, and right dorsal and ventral regions of middle frontal gyri, to a greater extent than do hits.

Functional data: parametric modulations

Parametric modulation analyses reveal regions that show an increasing (or decreasing) response as set size increases (or decreases). Such a pattern indicates that the regions contribute not only to true or false memory for items related to those studied previously (as shown with the random effects analyses), but show a graded response as set size increases.

Modulation with set size for false recognition

Activations associated with false alarms to increasing (Table 2A) or decreasing (Table 2B) set sizes are displayed in Fig. 3. For false recognition, as set size increased, activity increased in visual regions (fusiform and calcarine gyri) and the hippocampus.

Modulation with set size for true recognition

Activations associated with hits to increasing (Table 2C) or decreasing set sizes (Table 2D) are displayed in Fig. 3. In contrast to the pattern for false recognition, as set size increased for true recognition, activity decreased in visual regions (fusiform and middle occipital). In contrast, angular gyrus exhibits greater deactivation as set size decreases.

As noted in Table 2, many of the regions emerging in the parametric modulation analyses also reached significance in the random effects analyses, suggesting that there is consistency in the results from the two analyses.

Differences in modulation across true and false recognition

Because parametric modulation responses were analyzed separately for true and false recognition, we directly compared the patterns of responses across the two types of memory using paired samples *t*-tests in SPM, as described in the **Functional MRI analyses** section. Mean betas (see Table 3) reflect the average estimated slope of the regression line extracted from each condition; we then tested whether the betas differed significantly from zero. Regions in Table 3A exhibited positive modulations to false alarms (i.e., a stronger response as the amount of gist increases) and negative modulations (i.e., a stronger response as the amount of gist decreases) to hits (Table 3A), and included right hippocampus, parahippocampal gyrus, bilateral fusiform gyri, and occipital regions. Table 3B indicates regions showing positive modulations to hits and negative modulations to false alarms, and included temporal, frontal, and occipital regions.

Discussion

The findings of the present study are generally consistent with previous literature (e.g., Garoff-Eaton, et al., 2006; Slotnick and Schacter, 2004) in underscoring the extensive overlap in the sensory and reconstructive processes that support true and false recognition (Johnson et al., 1993; Mitchell and Johnson, 2009; Schacter and

Table 2

The table lists regions modulating hit or false alarm activity that surpass a voxel threshold of $p > .01$ and a cluster extent threshold > 44 to achieve an overall correction of $p < .05$. The top peak of each cluster is listed, and an asterisk (*) denotes regions that also emerge in the random effects analysis of large vs. small set sizes.

Region	BA	MNI coordinates of activation peak (x, y, z)			# of voxels	t value
A. False alarms: positive modulations (Lrg > Med > Small)						
*R fusiform	37	36	-38	-26	111	6.03
*R hippocampus	N/A	38	-10	-18	73	4.34
*L lingual/calcarine gyrus	17	2	-64	10	77	4.13
*R calcarine	17	16	-70	12	134	3.70
B. False alarms: negative modulations (Small > Med > Lrg)						
*R precentral	4	34	-24	58	97	4.80
*R caudate	25	14	24	0	113	4.04
*L superior parietal	7	-26	-48	58	53	4.03
C. Hits: positive modulations (Lrg > Med > Small)						
*R angular gyrus	39/40	58	-56	30	198	4.95
*L supramarginal gyrus	40/39	-60	-54	30	161	4.58
*L middle temporal/occipital	37/19	-54	-76	12	80	3.62
L inferior parietal	40/39	-48	-58	50	54	3.33
D. Hits: negative modulations (Small > Med > Lrg)						
*R inferior/middle occipital	19/18	38	-90	-2	333	5.39
*R middle occipital	19	32	-76	32	156	5.09
*L middle occipital	18	-30	-92	10	134	4.66
*R fusiform	37	38	-60	-14	205	4.27
*L putamen	N/A	-16	12	0	58	5.07
*L putamen	N/A	-28	-12	12	62	4.17
*L putamen	N/A	-24	14	12	53	4.12
*R cerebellum	37	36	-52	-24	50	4.44
*N/A	N/A	-18	40	14	49	4.37
*Vermis	N/A	0	-42	-16	61	3.86
*L midcingulate/paracentral	4	-14	-32	52	57	3.82
N/A	47	28	38	4	68	3.27

Addis, 2007; Schacter and Slotnick, 2004; Schacter, et al., in press). The results are novel because they provide information regarding the response of the hippocampus and other regions across levels of relatedness and degree of gist. While previous studies of true and false recognition treated relatedness in an all-or-none fashion, the present study investigated true and false recognition as a function of the degree of relatedness, finding that many of the relevant processes and corresponding neural regions respond in a graded fashion. As the gist representation is strengthened for false memories, the hippocampus and visual regions respond to a greater extent, indicating roles for constructive and perceptual processes as the degree of gist increases for false memory. This finding is consistent with behavioral studies of gist-based false recognition (Koutstaal and Schacter, 1997) in suggesting that the processes underlying false recognition are engaged to a greater extent as more related information has been encountered. For true memories, however, multimodal association regions track increasing amount of gist, perhaps playing a role in integrating sensory information and representing the outputs of retrieval processes in order to inform decision processes (Wagner et al., 2005), but perceptual regions respond to the individuation of objects (as gist decreases). Throughout the remainder of the discussion, we consider these component processes and their contribution to true and false recognition as gist strength increases and decreases.

The results implicate a number of processes as shared across true and false recognition, consistent with prior work (Garoff-Eaton, et al., 2006). For the comparison of true vs. false recognition, regardless of the degree of gist, the engagement of visual regions, including occipital cortex and fusiform gyrus, suggests that perceptual processes are invoked as gist information increases for both true and false memories. This activity could reflect the shared visual features across exemplars in a set (e.g., the shape of a chair). The activity of the

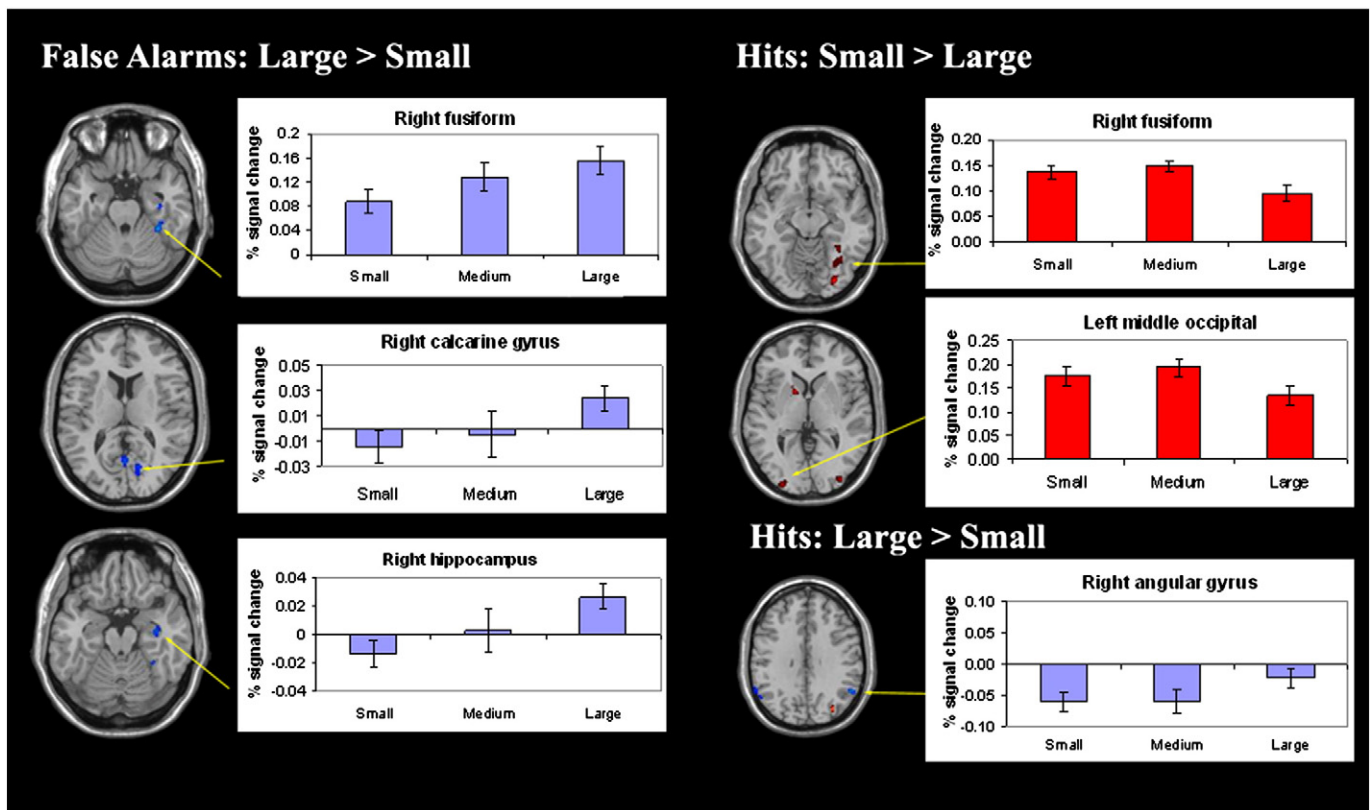


Fig. 3. Parametric modulations across set sizes. For false memories (left panel), fusiform (peak: 36, –38, –26) and calcarine (16, –70, 12) gyri activations reflect the role of visual regions in perceiving and recollecting prototypical features, and hippocampal (38, –10, –18) activations reflect the constructive nature of memory processes. These regions are more engaged for stronger gist representations. For true memories (right panel), the visual regions (fusiform [38, –60, –14] and left middle occipital [–30, –92, 10] gyri) are more engaged for more distinct items, reflecting a response to the visual properties of individual items, whereas angular gyrus (58, –56, 30) activations increase with set size, suggesting greater abstraction of conceptual and multimodal information as gist increases. Note that scales are unique to each graph/region.

hippocampus, in turn, could underlie the combination of perceptual, as well as semantic, features through the kinds of constructive memory processes discussed earlier (e.g., Schacter and Addis, 2007).

In terms of overall differences across true and false memories regardless of the level of gist, recognition is associated with the greater recruitment of visual, parahippocampal, posterior parietal, and temporal regions, likely reflecting increased access to sensory information for true than false recognition (e.g., Cabeza et al., 2001; Slotnick and Schacter, 2004). In contrast, false recognition is associated with greater engagement of frontal cortex, including anterior cingulate and right middle frontal gyrus, likely reflecting increased monitoring of retrieved memories (e.g., Cabeza et al., 2001; Schacter et al., 1996; Schacter et al., 1997; Slotnick and Schacter, 2004). These findings fit generally with the source monitoring framework in that true memories tend, on average, to have more perceptual details than false memories (Johnson, et al., 1993; Mitchell and Johnson, 2009). A number of results are consistent with this framework (e.g., Cabeza, et al., 2001; Gonsalves and Paller, 2000; Hashtroudi et al., 1990; Schacter, et al., 1996; Schacter, et al., 1997; Slotnick and Schacter, 2004). In contrast, frontal areas are more active when more monitoring is required (see Mitchell and Johnson, 2009 for a review), suggesting that in the present study, category-related lures may have generated more uncertainty than old items, engaging more frontally-based monitoring processes.

For true recognition, multimodal association areas were more engaged as gist strength increased. Activity linearly increased in regions including supramarginal and inferior parietal cortex as the number of related items increased. These parietal regions play a role in integrating information across modalities, reflecting the number of modalities engaged by objects (Wagner, et al., 2005). The

involvement of these regions suggests that integration of information drawn from multiple senses, in conjunction with semantic and linguistic processes (e.g., Binder et al., 2009), plays important roles in supporting accurate memory as more items were studied. As gist strength decreased, visual regions (with distinct peaks from those that modulated false alarms) were engaged, suggesting individuation of exemplars based on their unique perceptual features. For example, regions involved in visual analysis of objects (e.g., fusiform and middle occipital gyri) may contribute to individuating objects. These regions linearly increase their activity for hits as the number of related items decrease, suggesting a response to the visual properties of specific exemplars.

In terms of false memory, participants committed more false alarms as gist strength increased with larger set sizes at encoding, consistent with previous research (Koutstaal and Schacter, 1997). For false alarms, several regions were also associated with correct recognition (despite differences in peaks between true and false recognition), such as the fusiform gyrus, calcarine gyrus, and hippocampus, exhibited a larger response as the set size increased. Prior work has noted a role for these regions in accurate memory and false recognition of related items (Cabeza, et al., 2001; Garoff-Eaton, et al., 2006; Schacter and Slotnick, 2004; Slotnick and Schacter, 2004), and we will further consider the contributions of the hippocampus and visual regions.

Specifically, the hippocampus exhibited linear increases as encoding set size increased. Our data are consistent with the idea that the hippocampus contributes to constructive memory in that it flexibly binds elements together in memory, sometimes resulting in false memories through erroneous recombinations (Cabeza, et al., 2001; Giovanello, et al., 2010; Schacter and Addis, 2007; Schacter and

Table 3

Regions showing different patterns of modulation across false alarms (FA) and hits. An overall correction of $p < .05$ is achieved using a voxel extent threshold of $p < .01$ and a cluster extent threshold of 44 voxels. The mean beta indicates the direction of the effect for hits and false alarms. * indicates whether the positive or negative mean beta for hits or FA differs significantly from 0 at a threshold of $p < .01$.

Region	BA	MNI coordinates of activation peak (x, y, z)			# of voxels	t value	Mean beta	
							Hits	FA
<i>A. False alarms > hits</i>								
R fusiform	37	34	-38	-26	1044	5.51	-.81	1.67*
R inferior occipital	19	44	-74	-12		5.34	-1.25*	1.57*
R cerebellum		34	-52	-24		5.30	-1.96*	.55
R fusiform/ parahippocampal gyrus	20	38	-16	-24	87	3.92	-1.22*	.92*
R hippocampus	N/A	42	-18	-16		3.26	-.78*	.53
L fusiform	19	-34	-70	-16	45	3.33	-.98	.51
L middle occipital	18	-28	-92	8	70	3.67	-1.86*	.83
		-28	-86	2		3.10	-.59	.71
R mid-cingulate	23	10	-10	32	87	4.44	-.77*	.59
Midline	N/A	-4	-20	-8	48	4.06	-.62	.69*
Ventricle	N/A	-14	-34	26	108	6.69	-.56	.68*
White matter	N/A	-24	-28	24		3.58	-.51	.47
<i>B. Hits > false alarms</i>								
L sup temporal pole	38	-44	20	-12	56	5.07	.77	-1.12*
L inferior orbitofrontal	47	-34	22	-12		3.21	.61	-.82
L superior frontal	10	-16	60	20	47	4.55	.32	-.59
R precentral	4	34	-24	56	63	4.28	.57	-.86*
R superior temporal	22	64	-58	24	76	4.18	.91	-1.14*
R angular	40/39	64	-54	34		3.41	.84	-.72
L middle occipital	37	-44	-66	6	69	4.18	.97	-.96*
L middle temporal	37/19	-54	-72	14		3.18	1.75*	-1.11
R middle temporal	39	46	-66	20	64	4.06	.76*	-.61
	37	52	-68	14		3.00	.91	-.49

Slotnick, 2004; Slotnick and Schacter, 2004). The direct comparison of modulations for true and false memory suggests that the differential hippocampal activity tended to be the strongest for hits with small set size vs. large set size. The complex pattern of hippocampal engagement suggests reasons why previous studies have found an inconsistent role for the hippocampus in true and false memories. The engagement of the hippocampus could reflect the amount of information that needs to be bound together (captured here, but not in all previous studies, as the modulation across set size) as well as the content of that information (illustrated here through the differential effects of set size across hits and false alarms). For example, the hippocampus could be engaged by binding together more veridical details for small set size hits but more erroneous and gist-based details in the case of large set size false alarms.

An alternative explanation is that hippocampal activity reflects pattern separation mechanisms such as the greater demands of assigning a distinct representation to the lure stimulus (Bakker et al., 2008; Norman and O'Reilly, 2003). This interpretation fits well with prior data in that pattern separation, represented by engagement of areas in the CA3/dentate gyrus field of the hippocampus, is most required for novel information and similar lures, but less engaged for repeated information (Bakker, et al., 2008; Lacy et al., 2011). While it may seem counterintuitive to find a region more engaged for high amounts of gist for false, but not true, memory, it could reflect different contributions of pattern separation mechanisms to true vs. false memory. For example, greater hippocampal activity could be required in the case of true memories when few, specific veridical details were encoded (e.g., when distinct, nonoverlapping representations to novel stimuli were rapidly assigned) and

in the case of false memory, when prototypical features are falsely recalled (e.g., overlapping representations were assigned to similar stimuli, and high levels of overlap occur). In contrast, true memory influenced by high amounts of gist could depend on pattern completion mechanisms. Further work is needed because the pattern here is somewhat at odds with Norman and O'Reilly (2003) model (simulation 3), and because the contribution and failures of pattern separation and pattern completion mechanisms are not well understood across true and false memories sharing large amounts of gist.

Our results for greater involvement of visual regions as gist-based false recognition increases are consistent with the idea that people commit gist-based memory errors in this task due to mistakenly retrieving perceptual information (e.g., Mitchell and Johnson, 2009; Schacter and Slotnick, 2004; Schacter, et al., in press). For false memories, retrieving perceptual information is erroneous, leading individuals to falsely endorse novel items as familiar, based on their subjective experience (Johnson, et al., 1993). Interestingly, distinct visual regions are implicated for false alarms to new exemplars from large sets of encoded items. Specifically, activity in regions of the fusiform gyrus and calcarine cortex linearly increase as a function of increasing encoded set size. While prior work showed that early visual regions tend to be engaged by true more than false memories but respond to both types of memories to some degree (Slotnick and Schacter, 2004), our results suggest that these regions can be modulated by the strength of the gist component. Notably, these visual regions responded to the amount of gist information for both true and false recognition. This finding suggests that visual processes can benefit recognition by individuating small set exemplars but can also impair memory, perhaps by accentuating the prototypical visual features (e.g., the overall shape of a glove) common to novel and studied exemplars from large sets. The perceptual processes engaged could reflect a response to specific features viewed previously (e.g., the straight-back of a studied chair) or the strength of a pictorial representation of a prototypical, but unstudied, member of the relevant category. The potential importance of prototypical features is consistent with Koutstaal et al. (2003), who observed that semantic knowledge contributes to gist-based memory; in the present study, semantic knowledge of typical shape and perceptual properties could account for false memories. The potential importance of global shape information, likely to be shared across exemplars in a set, is also consistent with some work in perception (Sampanes et al., 2008). Future research is needed to better distinguish the component visual processes implicated in gist-based memory, particularly to address the question of whether the visual response is to specific features vs. prototypes and, importantly, whether the same vs. different visual processes engage the overlapping neural regions for true and false memory in this task.

In contrast to previous work (e.g., Dennis, et al., 2007; Dennis, et al., 2008; Garoff-Eaton, et al., 2007; Kim and Cabeza, 2007), temporal activations and the left inferior frontal activity that purportedly reflect semantic processing do not emerge in the present study as regions that respond as the amount of gist increases across true and false recognition. While it is possible that this divergence from prior literature reflects the fact that semantic processes per se do not contribute to false memories in a graded fashion (note that semantic processes could also engage perceptual processes, as discussed above), other explanations may account for this pattern. The engagement of temporal and inferior frontal regions in previous studies could reflect specific processes evoked by associative memories that have been implicated in DRM paradigms, rather than gist memory in the present paradigm. It is also possible that our use of pictures of nameable objects that tended to share both rich perceptual and conceptual information across exemplars evoked distinct processes than previous studies of related false recognition that used perceptually similar abstract shapes (e.g., Garoff-Eaton, et al., 2006; Slotnick and Schacter, 2004) or semantically related words (Dennis et al., 2007,

2008; Garoff-Eaton, et al., 2007; Kim and Cabeza, 2007; Schacter et al., 1996, 1997). We also implicate additional regions in false recognition that have not been emphasized in prior studies, with inferior parietal and middle and inferior prefrontal activity possibly tracking the amount of recollected information (Wagner, et al., 2005) and the engagement of monitoring processes (e.g., was this perceptual feature actually perceived, or was it invoked by studying other exemplars?) during the memory search (Schacter and Slotnick, 2004). Further work is needed to directly compare the activity evoked in this study with other false memory paradigms.

While the present study investigated the veridical and erroneous memory processes during the retrieval stage, it would also be of interest to consider the processes during encoding that lead to the formation of true and false memories, as well as those that contribute to the formation of gist representations. The majority of previous studies (e.g., Kim and Cabeza, 2007; Okado and Stark, 2003; Dennis et al., 2007; Dennis et al., 2008; Garoff-Eaton et al., 2007; Slotnick and Schacter, 2004; Kensinger and Schacter, 2005; Kensinger and Schacter, 2006) investigated the stages of memory separately, but the joint investigation of encoding and retrieval would allow for investigation into the interplay of processes across these stages. For example, it is possible that similar perceptual processes are invoked during encoding and retrieval, which would allow for a direct test of the sensory reactivation hypothesis such that robust engagement of sensory regions during encoding is mirrored by reactivation of the same regions at recognition. However, it is also possible that robust activation of later perceptual regions during encoding could support the formation of false memories due to shared higher-level features across many exemplars in a category.

In conclusion, our results converge with prior work in that false memories engage many of the same regions as true memory. However, we also find that perceptual and constructive, or binding, processes contribute to the degree of relatedness increases, thereby providing initial insights into the processes that are involved in the building of false memories that are based on the encoding and retrieval of gist information and further advancing our understanding of how erroneous memories can arise from otherwise adaptive cognitive processes.

Supplementary materials related to this article can be found online at doi:10.1016/j.neuroimage.2011.11.078.

Acknowledgments

The authors gratefully acknowledge support from the National Institutes of Health, grants NIMH R01 MH60941 (to DLS) and NIA F32 AG026920 (to AHG). Portions of this research were conducted when A.H.G. was a fellow of the American Federation for Aging Research. The Athinoula A. Martinos Center for Biomedical Imaging is supported by the National Center for Research Resources (grant P41 RR14075) and by the MIND Institute. We thank Elizabeth Kensinger and Rachel Garoff Eaton for helpful discussions, and Donna Rose Addis for generously sharing her expertise and insights.

References

- Abe, N., Okuda, J., Suzuki, M., Sasaki, H., Matsuda, T., Mori, E., et al., 2008. Neural correlates of true memory, false memory, and deception. *Cereb. Cortex* 18, 2811–2819.
- Addis, D.R., Schacter, D.L., 2008. Constructive episodic simulation: temporal distance and detail of past and future events modulate hippocampal engagement. *Hippocampus* 18, 227–237.
- Addis, D.R., Wong, A.T., Schacter, D.L., 2007. Remembering the past and imagining the future: common and distinct neural substrates during event construction and elaboration. *Neuropsychologia* 45, 1363–1377.
- Addis, D.R., Pan, L., Vu, M.A., Laiser, N., Schacter, D.L., 2009. Constructive episodic simulation of the future and the past: distinct subsystems of a core brain network mediate imagining and remembering. *Neuropsychologia* 47, 2222–2238.
- Addis, D.R., Cheng, T., Roberts, R., Schacter, D.L., 2011. Hippocampal contributions to the episodic simulation of specific and general future events. *Hippocampus* 21, 1045–1052.
- Bakker, A., Kirwan, C.B., Miller, M., Stark, C.E.L., 2008. Pattern separation in the human hippocampal CA3 and dentate gyrus. *Science* 319, 1640–1642.
- Bartlett, F.C., 1932. *Remembering*. Cambridge University Press, Cambridge.
- Binder, J.R., Desai, R.H., Graves, W.W., Conant, L.L., 2009. Where is the semantic system? A critical review and meta-analysis of 120 functional neuroimaging studies. *Cereb. Cortex* 19, 2767–2796.
- Brainerd, C.J., Reyna, V.F., 2002. Fuzzy trace theory and false memory. *Curr. Dir. Psychol. Sci.* 11, 164–169.
- Brett, M., Anton, J.-L., Valabregue, R., Poline, J.-B., 2002. Region of interest analysis using an SPM toolbox. Paper presented at the 8th International Conference on Functional Mapping of the Human Brain, Sendai, Japan.
- Buckner, R.L., 2010. The role of the hippocampus in prediction and imagination. *Annu. Rev. Psychol.* 61, 27–48.
- Cabeza, R., Rao, S.M., Wagner, A.D., Mayer, A.R., Schacter, D.L., 2001. Can medial temporal lobe regions distinguish true from false? An event-related functional MRI study of veridical and illusory recognition memory. *Proc. Natl. Acad. Sci. U. S. A.* 98, 4805–4810.
- Dale, A.M., 1999. Optimal experimental design for event-related fMRI. *Hum. Brain Mapp.* 8, 109–114.
- Deese, J., 1959. Influence of inter-item associative strength upon immediate free recall. *Psychol. Rep.* 5, 305–312.
- Dennis, N.A., Kim, H., Cabeza, R., 2007. Effects of aging on true and false memory formation: an fMRI study. *Neuropsychologia* 45, 3157–3166.
- Dennis, N.A., Hayes, S.M., Prince, S.E., Madden, D.J., Huettel, S.A., Cabeza, R., 2008. Effects of aging on the neural correlates of successful item and source memory encoding. *J. Exp. Psychol. Learn. Mem. Cogn.* 34, 791–808.
- Fisher, R.A., 1950. 11th ed. *Statistical Methods for Research Workers*, vol. 5. Hafner, New York.
- Gallate, J., Chi, R., Ellwood, S., Snyder, A., 2009. Reducing false memories by magnetic pulse stimulation. *Neurosci. Lett.* 449, 151–154.
- Gallo, D.A., 2006. *Associative Illusions of Memory: False Memory Research in DRM and Related Tasks*. Psychology Press, New York.
- Gallo, D.A., 2010. False memories and fantastic beliefs: 15 years of the DRM illusion. *Mem. Cogn.* 38, 833–848.
- Garoff-Eaton, R.J., Slotnick, S.D., Schacter, D.L., 2006. Not all false memories are created equal: the neural basis of false recognition. *Cereb. Cortex* 16, 1645–1652.
- Garoff-Eaton, R.J., Kensinger, E.A., Schacter, D.L., 2007. The neural correlates of conceptual and perceptual false recognition. *Learn. Mem.* 14, 684–692.
- Giovanello, K.S., Kensinger, E.A., Wong, A.T., Schacter, D.L., 2010. Age-related neural changes during memory conjunction errors. *J. Cogn. Neurosci.* 22, 1348–1361.
- Gonsalves, B., Paller, K.A., 2000. Neural events that underlie remembering something that never happened. *Nat. Neurosci.* 3, 1316–1321.
- Guerin, S.A., Robbins, C.R., Gilmore, A.W., Schacter, D.L. in press. Retrieval failure contributes to gist-based false recognition. *Journal of Memory and Language*.
- Hashtroudi, S., Chrosniak, L.D., Johnson, M.K., 1990. Aging and qualitative characteristics of memories for perceived and imagined complex events. *Psychol. Aging* 5, 119–126.
- Hassabis, D., Kumaran, D., Vann, S.D., Maguire, E.A., 2007. Patients with hippocampal amnesia cannot imagine new experiences. *Proc. Natl. Acad. Sci. U. S. A.* 104, 1726–1731.
- Johnson, M.K., Hashtroudi, S., Lindsay, D.S., 1993. Source monitoring. *Psychol. Bull.* 114, 3–28.
- Johnson, M.K., Nolde, S.F., Mather, M., Kounios, J., Schacter, D.L., Curran, T., 1997. The similarity of brain activity associated with true and false recognition depends on the test format. *Psychol. Sci.* 8, 250–251.
- Kensinger, E.A., Schacter, D.L., 2005. Retrieving accurate and distorted memories: neuroimaging evidence for effects of emotion. *NeuroImage* 27, 167–177.
- Kensinger, E.A., Schacter, D.L., 2006. Neural processes underlying memory attribution on a reality-monitoring task. *Cereb. Cortex* 16, 1126–1133.
- Kim, H., Cabeza, R., 2007. Differential contributions of prefrontal, medial temporal, and sensory-perceptual regions to true and false memory formation. *Cereb. Cortex* 17, 2143–2150.
- Koutstaal, W., 2003. Older adults encode—but do not always use—perceptual details: intentional versus unintentional effects of detail on memory judgments. *Psychol. Sci.* 14, 189–193.
- Koutstaal, W., Cavendish, M., 2006. Using what we know: consequences of intentionally retrieving gist versus item-specific information. *J. Exp. Psychol. Learn. Mem. Cogn.* 32, 778–791.
- Koutstaal, W., Schacter, D.L., 1997. Gist-based false recognition of pictures in older and younger adults. *J. Mem. Lang.* 555–583.
- Koutstaal, W., Schacter, D.L., Galluccio, L., Stofer, K.A., 1999. Reducing gist-based false recognition in older adults: encoding and retrieval manipulations. *Psychol. Aging* 14, 220–237.
- Koutstaal, W., Reddy, C., Jackson, E.M., Prince, S., Cendan, D.L., Schacter, D.L., 2003. False recognition of abstract versus common objects in older and younger adults: testing the semantic categorization account. *J. Exp. Psychol. Learn. Mem. Cogn.* 29, 499–510.
- Lacy, J.W., Yassa, M.A., Stark, S.M., Muftuler, L.T., Stark, C.E.L., 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, 15–18.
- Loftus, E.F., 1979. Malleability of human memory. *Am. Sci.* 67, 312–320.
- Martin, V.C., Schacter, D.L., Corballis, M., Addis, D.R., 2011. A role for the hippocampus in encoding simulations of future events. *Proc. Natl. Acad. Sci. U. S. A.* 108, 13858–13863.
- Mather, M., Henkel, L.A., Johnson, M.K., 1997. Evaluating characteristics of false memories: remember/know judgments and memory characteristics questionnaire compared. *Mem. Cogn.* 25, 826–837.

- Mitchell, K.J., Johnson, M.K., 2009. Source monitoring 15 years later: what have we learned from fMRI about the neural mechanisms of source memory? *Psychol. Bull.* 135, 638–677.
- Mitchell, K.J., Johnson, M.K., Raye, C.L., D'Esposito, M., 2000. fMRI evidence of age-related hippocampal dysfunction in feature binding in working memory. *Cogn. Brain Res.* 10, 197–206.
- Neisser, U., 1967. *Cognitive Psychology*. Appleton-Century-Crofts, New York.
- Newman, E.J., Lindsay, D.S., 2009. False memories: what the hell are they for? *Appl. Cogn. Psychol.* 23, 1105–1121.
- Norman, K.A., O'Reilly, R.C., 2003. Modeling hippocampal and neocortical contributions to recognition memory: a complementary-learning-systems approach. *Psychol. Rev.* 110, 611–646.
- Okado, Y., Stark, C., 2003. Neural processing associated with true and false memory retrieval. *Cogn. Affect. Behav. Neurosci.* 3, 323–334.
- Reyna, V.F., Brainerd, C.J., 1995. Fuzzy trace theory – an interim synthesis. *Learn. Individ. Differ.* 7, 1–75.
- Roediger III, H.L., McDermott, K.B., 1995. Creating false memories: remembering words not presented on lists. *J. Exp. Psychol. Learn. Mem. Cogn.* 21, 803–814.
- Roediger III, H.L., McDermott, K.B., 2000. Tricks of memory. *Curr. Dir. Psychol. Sci.* 9, 123–127.
- Rorden, C., Brett, M., 2000. Stereotaxic display of brain lesions. *Behav. Neurol.* 12, 191–200.
- Sampanes, A.C., Tseng, P., Bridgeman, B., 2008. The role of gist in scene recognition. *Vis. Res.* 48, 2275–2283.
- Schacter, D.L., 2001. *The Seven Sins of Memory: How the Mind Forgets and Remembers*. Houghton Mifflin, Boston.
- Schacter, D.L., Addis, D.R., 2007. The cognitive neuroscience of constructive memory: remembering the past and imagining the future. *Philos. Trans. R. Soc. B-Biol. Sci.* 362, 773–786.
- Schacter, D.L., Addis, D.R., 2009. On the nature of medial temporal lobe contributions to the constructive simulation of future events. *Philos. Trans. R. Soc. B-Biol. Sci.* 364, 1245–1253.
- Schacter, D.L., Slotnick, S.D., 2004. The cognitive neuroscience of memory distortion. *Neuron* 44, 149–160.
- Schacter, D.L., Reiman, E., Curran, T., Yun, L.S., Bandy, D., McDermott, K.B., et al., 1996. Neuroanatomical correlates of veridical and illusory recognition memory: evidence from positron emission tomography. *Neuron* 17, 267–274.
- Schacter, D.L., Buckner, R.L., Koutstaal, W., Dale, A.M., Rosen, B.R., 1997. Late onset of anterior prefrontal activity during true and false recognition: an event-related fMRI study. *NeuroImage* 6, 259–269.
- Schacter, D.L., Norman, K.A., Koutstaal, W., 1998. The cognitive neuroscience of constructive memory. *Annu. Rev. Psychol.* 49, 289–318.
- Schacter, D.L., Israel, L., Racine, C., 1999. Suppressing false recognition: the distinctiveness heuristic. *J. Mem. Lang.* 40, 1–24.
- Schacter, D.L., Guerin, S.A., St. Jacques, P.L., 2011. Memory distortion: an adaptive perspective. *Trends Cogn. Sci.* 15, 467–474.
- Schacter, D.L., Chamberlain, J., Gaesser, B., Gerlach, K.D. in press. Neuroimaging of true, false, and imaginary memories: findings and implications. In L. Nadel & W. Sinnott-Armstrong (Eds.), *Memory and Law: Perspectives from Cognitive Neuroscience*. New York: Oxford University Press.
- Slotnick, S.D., Schacter, D.L., 2004. A sensory signature that distinguishes true from false memories. *Nat. Neurosci.* 7, 664–672.
- Slotnick, S.D., Moo, L.R., Segal, J.B., Hart Jr., J., 2003. Distinct prefrontal cortex activity associated with item memory and source memory for visual shapes. *Cogn. Brain Res.* 17, 75–82.
- Talairach, J., Tournoux, P., 1988. *Co-planar Stereotaxic Atlas of the Human Brain*. (M. Rayport, Trans.) Thieme Medical Publishers, Inc., New York.
- Tun, P.A., Wingfield, A., Rosen, M.J., Blanchard, L., 1998. Response latencies for false memories: gist-based processes in normal aging. *Psychol. Aging* 13, 230–241.
- Tzourio-Mazoyer, N., Landeau, B., Papathanassiou, D., Crivello, F., Etard, O., Delcroix, N., et al., 2002. Automated anatomical labeling of activations in SPM using a macroscopic anatomical parcellation of the MNI MRI single-subject brain. *NeuroImage* 15, 273–289.
- Wagner, A.D., Shannon, B.J., Kahn, I., Buckner, R.L., 2005. Parietal lobe contributions to episodic memory retrieval. *Trends Cogn. Sci.* 9, 445–453.