Preserved verbal memory function in left medial temporal pathology involves reorganisation of function to right medial temporal lobe

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Abstract

The left hippocampus and related structures mediate verbal memory function. The mechanism underlying preserved verbal memory function in patients with left hippocampal damage is unknown. Temporal lobe epilepsy, a common disease, is frequently the consequence of a characteristic hippocampal pathology termed hippocampal sclerosis, which may also affect the amygdala. In this setting, mapping the sites of memory function is a vital component of planning for surgical treatment for epilepsy. Using event-related functional magnetic resonance imaging, we studied 24 right-handed nonamnesic patients with left hippocampal sclerosis and 12 normal controls, performing a verbal encoding task. The patients were subdivided into two groups according to presence or absence of additional left amygdala pathology. Analysis of the data employed a two-level random-effects design, examining the main effects of subsequent memory in each group, as well as the differences between the groups. Additional effects of emotionality of the remembered words were also examined. Verbal memory encoding involved activation of left hippocampus in normals, but was associated with reorganisation to right hippocampus and parahippocampal gyrus in the patients. The additional presence of left amygdala sclerosis resulted in reorganisation for encoding of emotional verbal material to right amygdala. Retained verbal memory function in the presence of left medial temporal lobe pathology is mediated by recruitment of a parallel system in the right hemisphere consistent with adaptive functional reorganisation. The findings indicate a high degree of plasticity in medial temporal lobe structures.

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Introduction

The human hippocampus and related medial temporal lobe (MTL) structures are crucial to long-term episodic memory function (Squire, 1992). Studies of subjects with injury to this area reveal a characteristic syndrome of amnesia (Squire and Zola-Morgan, 1991). The role of the hippocampus has been confirmed and extended by functional imaging in normal subjects which reveal hippocampal activity in long-term memory tasks (Lepage et al., 1998; Cabeza and Nyberg, 2000). A dissociation between left hippocampus, subserving verbal memory, and right hippocampus, mediating visual memory, is evident from studies of patients (Smith and Milner, 1981; Frisk and Milner, 1990) and functional neuroimaging in normal subjects (Kelley et al., 1998).

The hippocampus is the site of a common pathology, hippocampal sclerosis (HS), which frequently underlies a common disease, temporal lobe epilepsy (TLE). Although patients with TLE may have memory impairment, amnesia is unusual (Helmstaedter and Kurthen, 2001). HS is detected in vivo using magnetic resonance imaging (MRI) (Duncan, 1997) and its hallmarks are reduced hippocampal volume and increased T2 signal, with other medial temporal structures and more remote brain regions generally showing no pathology (Sisodiya et al., 1997). Although bilateral HS is found in a minority, standardised imaging criteria used in this study allow unilateral HS to be identified (Woermann et al., 1998). Therefore, subjects with HS provide a unique opportunity to study long-term consequences of isolated unilateral hippocampal injury.
Given the role of left hippocampus in verbal memory function, the observation that verbal episodic memory is relatively intact in subjects with left HS suggests reorganisation of episodic memory processes to other brain loci. During evaluation for neurosurgical treatment of epilepsy, TLE patients may undergo a procedure in which first one hemisphere, then the other, is anaesthetised by a short-acting barbiturate injected into the internal carotid artery (Wada test), allowing the memory capacity of each unanaesthetised hemisphere to be tested in isolation. Usually, memory is superior in the hemisphere unaffected by HS. Bilateral temporal lobe resection in TLE patients produces devastating amnesia, as in the famous amnesic patient HM (Scoville and Milner, 1957). This implies that in unilateral HS, memory function is reallocated from the hippocampus affected by HS to the contralateral temporal lobe.

Anterior to the hippocampus is the amygdala, a region linked to emotional perception and memory (Dolan, 2002). A close functional relationship between amygdala and hippocampus is suggested by enhancement in free recall of emotionally significant relative to neutral material (Phelps et al., 1997). Patients with isolated amygdala damage show a specific impairment in recall of emotional stimuli (Adolphs et al., 1997). Functional imaging studies show that amygdala activity correlates with subsequent memory for emotional material (Hamann et al., 1999; Strange et al., 2000). In patients with HS, the amygdala may be affected by the same sclerotic pathology as the hippocampus, although the severity of hippocampal and amygdala sclerosis in a given patient may be uncorrelated (Hudson et al., 1993). As in the case of the hippocampus, amygdala pathology may be assessed using MRI by measuring T2 signal (Van Paesschen et al., 1996; Bartlett et al., 2002).

Functional imaging studies have frequently sought evidence of the functional adaptation of the brain to the presence of a lesion in the brain or peripheral nervous system. Few instances have emerged in which good task performance is associated with a pattern of brain activity clearly different from that in normal subjects. Following stroke, clinical recovery is related to functional recovery within the damaged area itself, with reorganisation of function to remote brain regions (such as the opposite undamaged hemisphere) being associated with poor recovery (Warburton et al., 1999; Pineiro et al., 2001; Johansen-Berg et al., 2002). Reorganisation of brain function resulting in good performance has been demonstrated for sensory functions as evidenced in long-term deaf (Neville et al., 1998) or blind (Sadato et al., 1996) adults. Reorganisation of language to right frontal regions has been reported in recovered aphasics with left frontal lesions (Blasi et al., 2002), in addition to reorganisation of motor function to the ipsilateral hemisphere in adults with congenital hemiparesis (Staudt et al., 2002). The ability of regions outside the left MTL to subserve residual verbal memory functions has not been demonstrated.

In this study, we describe a functional MRI (fMRI) study of verbal encoding in subjects with an MRI-based diagnosis of unilateral left HS, compared to normal controls. We hypothesised that pathology in left medial temporal lobe structures would result in reallocation of verbal encoding functions to the right MTL. As functional neuroimaging studies of patients may be confounded by heterogeneity of sample and poor task performance compared to normals, we studied a homogeneous group of right-handed patients with identical pathology and compared encoding event types which showed the least variance between patients and normals.

**Materials and methods**

**Subjects**

We studied 12 right-handed normal volunteers and 24 right-handed patients recruited from the National Hospital for Neurology and Neurosurgery, London. All lived in southern England and were fluent English speakers. All patients had active temporal lobe epilepsy and had undergone MRI at 1.5 T (Horizon Echospeed, General Electric, Milwaukee, WI, USA) (Duncan et al., 1996; Duncan, 1997; Woermann et al., 1998), which had identified left HS and normal right hippocampus. The patients were divided into two groups based on left amygdala T2: the first group (HS) had normal left amygdala T2; the second group had normally elevated left amygdala T2 (HSAS) (Bartlett et al., 2002). The age range of the patient group was 18–54 (median 29) and of the normal control group was 19–48 (median 31); 10 patients and 8 normal subjects were male; all patients were on anticonvulsant medication. All patients had undergone neuropsychometry and had a verbal IQ >80 (range 80–124, median 102), IQ data were not obtained for the normal subjects. Approval for this study was obtained from the joint ethics committee of the hospital and the Institute of Neurology. All subjects gave written informed consent.

**Imaging**

Subjects were scanned at 2 T (Siemens Vision, Siemens, Erlangen, Germany), acquiring gradient-echo echo-planar T2*-weighted image volumes, providing blood oxygenation level-dependent contrast, with 33 contiguous slices covering the whole brain, voxel dimensions 3 × 3 × 3.67 mm, TE 40 ms, and TR 2.5 s. SPM99 was used for image analysis (Friston et al., 1995). The images were realigned and corrected for slice timing differences. All of the images from each subject were then transformed to the standard anatomical volume and smoothed with an 8-mm kernel.

**Psychological task**

During scanning subjects were presented 255 single words previously used elsewhere (Otten et al., 2001), including 36 emotionally aversive words (e.g., “cancer,” ...
rape,” “terrorist”) (Strange et al., 2000), 1 every 4.5 s. Subjects were instructed to indicate with a single right-hand button press whether the word indicated a living or nonliving entity. Ninety minutes after scanning, subjects performed a surprise recognition memory test outside fMRI scanning (Fig. 1). At recognition, single words were presented every 4.5 s in a manner identical to that used in scanning: 255 words presented during scanning, randomly mixed with 170 neutral foils and 36 emotional foils. For each stimulus, subjects were asked to indicate with a right-hand button press if they could remember seeing the word on the screen during scanning (R response), if the word seemed familiar (K response), or if the word was new (N response) (Tulving, 1985). The encoding stimuli were then conditionalised according to these recognition responses; seven event types were identified: correct R responses for neutral stimuli (Rn), correct K responses to neutral stimuli (Kn), incorrect N responses to neutral stimuli (forgotten, Fn), and similarly for emotional stimuli (Re, Ke, Fe). A single category for all encoding task errors or missed responses was included, hence in total seven event types were indicated. To calculate recognition accuracy for each event type (Rn, Kn, Re, Ke), stimuli seen in the recognition test were classified as correct identifications (hits) and incorrect responses to foils (false alarms). Recognition accuracy was calculated for each of these four event types as (hit rate) − (false alarm rate). To calculate the recognition accuracy for N responses, stimuli in the recognition test were similarly classified as correct rejections and misses; recognition accuracy for N responses was calculated as (miss rate) − (correct rejection rate), for neutral and emotional stimuli separately. The highest possible score was +1, indicating perfect identification of stimuli present at encoding; the lowest score, −1, indicated perfect identification of foils.

Imaging analysis

To test for subsequent memory effects, imaging data were analysed with a two-level random-effects analysis employing an event-related design (Friston et al., 1998). At the first level, trial-specific responses were modelled for each subject by convolving a delta function with the canonical haemodynamic response function (HRF) to create regressors of interest, one regressor for each of the seven event types described above. Each subject’s movement parameters were included as confounds. A high-pass filter of 120 s was used and data were scaled for global activity. Parameter estimates pertaining to the height of the HRF for each regressor were calculated at each voxel. Contrasts of parameter estimates were calculated to produce four “contrast images” for each subject (Rn minus Kn, Re minus Ke, main effect of R minus K, interaction between R minus K and emotion). These images were used for the second-level analyses.

At the second level, one-sample T tests were used to examine effects within groups, and two-sample T tests used to examine between-group effects. We chose p < 0.05 corrected (within the small volume of the MTL using an image mask) as the threshold for significance. We report all MTL activations that survived this threshold and report descriptively MTL regions showing uncorrected significance where we replicate previous findings or showing a nonsignificant trend where this contributes to interpretation of the data.

Results

Left and right hippocampal volumes were significantly different in the patients: mean left hippocampal volume 1834 mm³, mean right hippocampal volume 2797 mm³.
larger left hippocampi than HS patients (HS group left-tailed). There was a strong trend for HSAS patients to have versus 94.3 ms in HSAS group; (mean relaxation time 88.2 ms in HS group (HSAS) had abnormally elevated left amygdala T2 (Bartlett et al., 2002) (mean relaxation time 88.2 ms in HS group (HS) had normal left amygdala T2; the second group divided into two groups based on left amygdala T2: the

significant effect of group for either R or K responses (Fig. 1). Repeated-measures ANOVA with emotionality as a within-subjects factor revealed no significant effect of group (left hippocampus in normals compared to patients. Both hippocampi showed a two-way interaction between subsequent memory and emotionality, collapsed across group at p < 0.05 corrected (Figs. 3b and 3c). This effect was characterised by greater activity to emotional compared to neutral subsequently remembered stimuli. There were no regions showing greater activity to neutral items than to emotional items. Finally, the three-way interaction between memory, group (HS versus HSAS), and emotionality showed effects in an anterolateral region close to right amygdala and at the right amygdalohippocampal junction at p < 0.05 corrected; both regions were characterised by relatively greater activity for emotional items than for neutral items in HSAS patients compared with HS patients (Fig. 4). There were no other significant three-way interactions.

In a subsequent analysis, to test specifically for hemispheric lateralisation, the peak voxel values for the main effect of memory (R minus K) in left (−32, −18, −18) and right hippocampus (26, −14, −18) for all subjects were entered into a repeated-measures ANOVA with “hemisphere” (left or right) as a within-subjects effect and “group” (normals or patients) as a between-subjects effect. There was a significant interaction (p = 0.046) between hemisphere and group, explained by less activity in the right hippocampus in normals compared to patients.

Table 1
Activation peaks for subsequent memory effects in medial temporal lobe

<table>
<thead>
<tr>
<th>Contrast</th>
<th>Fig.</th>
<th>MNI coordinates</th>
<th>Z score</th>
<th>Region</th>
</tr>
</thead>
<tbody>
<tr>
<td>Main effect of memory, normal controls</td>
<td>2a</td>
<td>−32 −18 −18</td>
<td>2.35</td>
<td>Left hippocampus</td>
</tr>
<tr>
<td>Main effect of memory, patients</td>
<td>2b</td>
<td>18 −6 −20</td>
<td>4.29*</td>
<td>Right amygdala</td>
</tr>
<tr>
<td></td>
<td>2c</td>
<td>26 −14 −18</td>
<td>3.53*</td>
<td>Right hippocampus</td>
</tr>
<tr>
<td></td>
<td>2d</td>
<td>24 −28 −20</td>
<td>3.27*</td>
<td>Right parahippocampal gyrus</td>
</tr>
<tr>
<td></td>
<td>2e</td>
<td>−26 −24 −20</td>
<td>3.32*</td>
<td>Inferior to left hippocampus</td>
</tr>
<tr>
<td>Two-way interaction between memory and group (collapsed across emotionality)</td>
<td>3a</td>
<td>26 −20 −32</td>
<td>3.45*</td>
<td>Right parahippocampal gyrus</td>
</tr>
<tr>
<td></td>
<td>3b</td>
<td>−28 −22 −10</td>
<td>3.0*</td>
<td>Left hippocampus</td>
</tr>
<tr>
<td>Two-way interaction between memory and emotionality (collapsed across group)</td>
<td>3c</td>
<td>30 −18 −14</td>
<td>2.71</td>
<td>Right hippocampus</td>
</tr>
<tr>
<td>Three-way interaction between memory, emotionality, and group</td>
<td>4a</td>
<td>32 8 −22</td>
<td>3.77*</td>
<td>Right periamygdala</td>
</tr>
<tr>
<td></td>
<td>4b</td>
<td>26 −10 −14</td>
<td>3.0*</td>
<td>Right amygdala/hippocampus</td>
</tr>
</tbody>
</table>

* Regions significant at p < 0.05 corrected.

For each effect, the Montreal Neurological Institute (MNI) coordinates, Z score, anatomical location, and relevant figure are given.
**Discussion**

In nonamnesic patients with left MTL pathology we demonstrate reorganisation or reallocation of encoding processes to right MTL in two distinct ways: first, subjects with left hippocampal sclerosis (HS and HSAS groups combined) showed greater activity in right hippocampus and parahippocampal gyrus than normal subjects during successful encoding of words, compared to normal subjects. Second, subjects with left amygdala sclerosis (HSAS group) showed greater activity in the right amygdala compared to subjects without amygdala sclerosis (HS group) for successfully encoded emotional compared to neutral words. Hence, in these nonamnesic subjects with MTL pathology, verbal memory for both neutral and emotional items is associated with reallocation of encoding-related activity from left-sided MTL structures to equivalent structures in the right MTL.

Our study has a number of methodological strengths which enable a high degree of confidence in these findings. First, we examined a relatively large and highly homogeneous cohort of right-handed patients with clearly charac-

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Fig. 2. Main effects of R minus K. Significant regions (all voxels $p < 0.01$ in this and all subsequent figures) are superimposed onto an “average patient” T1 image, the voxel-by-voxel mean of the volumetrically normalised T1 images of all 24 patients. (a) Normal group left hippocampal peak. Although the statistical test pertains only to the main effect of memory (R minus K) and only to the normal subjects, parameter estimates are presented for all three groups and separately for neutral and emotional items in the bar chart. The $y$-axis represents activity in arbitrary units on this and all subsequent figures. Each bar represents the mean R minus K parameter estimate difference and the error bar ±1 SEM. Pale grey bars: neutral items. Dark grey bars: emotional items. (b) Combined patient groups, peak inferior to left hippocampus. Again, although the statistical test pertains only to the main effect of memory and only to the patients, parameter estimates are presented for all three groups and separately for neutral and emotional items in the bar chart. (c) Combined patient groups, right hippocampal peak. (d) Combined patient groups, right parahippocampal peak. (e) Combined patient groups, right amygdala peak.
terised MTL pathology. Second, we employed a psychological task which allowed good matching of task performance between normals and patients in conjunction with an event-related fMRI design which isolated the successful encoding processes of interest. Third, the encoding session during scanning included words with a highly emotional content, which led to robust engagement of MTL. Fourth, we employed a highly stringent, robust, and conservative statistical approach (multilevel modelling and “random effects”) which permits inference to be drawn about the population.

Fig. 3. Two-way interactions. (a) Interaction between main effect of memory and group (HS and HSAS combined versus normals): right parahippocampal peak. (b) Interaction between memory and emotionality (collapsed across all three groups): right hippocampus. (c) Interaction between memory and emotionality (collapsed across all three groups): left hippocampus.

Fig. 4. Three-way interactions. (a) Interaction between memory, emotionality, and group (HSAS versus HS): right periamygdalar region. (b) Interaction between memory, emotionality, and group (HSAS versus HS): right amygdalohippocampal junction.
from which subjects are drawn, rather than merely inference pertaining only to the subjects studied.

There have been several previous attempts to identify the site of residual memory function in subjects with left TLE. Subjects with left TLE showed less left MTL activity than subjects with right TLE during a block-design verbal task designed to identify regions engaged by semantic language subjects with right TLE during a block-design verbal task (Bellgowan et al., 1998); the MTL activity was designed to identify regions engaged by semantic language subjects with right TLE during a block-design verbal task to be superior to drug treatment for controlling seizures. Avoidance of postoperative memory impairment is a crucial objective in this setting (Baxendale, 1998). Although psychometric tests, structural imaging, and the Wada test may identify patients at special risk of postoperative amnesia, a need exists for functional mapping of residual memory abilities. The growth in understanding of the functional anatomy of memory in normal subjects has not been matched by knowledge concerning how the human brain compensates following injury to primary memory regions. Our study suggests that effective verbal memory encoding in subjects with left hippocampal and hippocampal plus amygdala damage is mediated by recruitment of parallel or redundant memory systems within the right MTL.

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**References**


