

Anterograde and retrograde amnesia in a person with bilateral fornix lesions following removal of a colloid cyst

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Abstract

AD, a 45-year-old man, presented with a severe and global anterograde amnesia following surgery for removal of a colloid cyst. Structural neuroimaging confirmed bilateral lesions to the fornix and a small lesion in the basal forebrain. Testing for remote episodic memory of autobiographical events, and for remote semantic memory of personal and public events, and of famous people, revealed that AD had a severe retrograde amnesia for autobiographical episodes that covered his entire lifetime, and a time-limited retrograde amnesia for semantic memory. Because the fornix and basal forebrain lesions disrupted major afferent and efferent pathways of the hippocampus, it was concluded that the integrity of the hippocampus and its projections are needed to retain and/or recover autobiographical memories no matter how old they are. By contrast, hippocampal contribution to semantic memory is time-limited. These findings were interpreted as consistent with Multiple Trace Theory, which holds that the hippocampal system is essential for recovering contextually rich memories no matter how old they are, but is not needed for recovering semantic memories.

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1. Introduction

It is widely accepted that bilateral damage to the hippocampus and related medial temporal lobe (MTL) structures, the perirhinal, entorhinal and parahippocampal cortices, leads to severe anterograde amnesia. However, there is considerable debate regarding the effects of damage to this region on remote memory. The debate turns on several points: (1) the location and extent of the lesion; (2) the temporal extent and severity of the deficit, and (3) the type of memory which is affected. With respect to the first two points, studies show that damage restricted to the MTL, which includes the hippocampus, leads to remote memory loss, the severity and extent being determined by the location and size of the lesion (Aggleton and Brown, 1999). Others argue that the

lesion must extend to extra-MTL structures to produce a temporally extensive retrograde amnesia (Bayley, Gold, Hopkins, & Squire, 2005; Bayley, Hopkins and Squire, 2003). With respect to the type of memory deficit in patients with damage limited to the MTL, there is an emerging consensus that memory for specific personal information (episodic memory) is severely affected, with memory for general knowledge and factual information about the world and oneself (semantic memory) being relatively spared (for review see Moscovitch, Rosenbaum et al., 2005). This evidence is consistent with Multiple Trace Theory which posits that the hippocampus and related MTL structures are always needed for retention and recovery of contextually rich information, which is the hallmark of episodic memory. On the other hand, memories that are independent of the context in which they were acquired, such as semantic memories, can be mediated through extra-hippocampal structures (Nadel & Moscovitch, 1997; Rosenbaum, Winocur, & Moscovitch, 2001).

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It has been difficult to evaluate the contribution of the hippocampus to remote memory because rarely, if ever, is damage restricted to this structure in humans. Furthermore, when it is, the hippocampal lesion is smaller than in patients whose lesions extend beyond this structure. In other words, we do not have a case where most of the hippocampus is destroyed without also encroaching on extra-hippocampal structures. We address these issues in a patient, AD, with bilateral lesions to the fornix, a major pathway linking the hippocampus to basal forebrain and diencephalic regions implicated in memory. The fornix is the primary efferent projection from the hippocampus to the mammillary bodies and anterior thalamic nuclei, and also contains afferent cholinergic tracts from the septal nuclei in the basal forebrain to the hippocampus. To a lesser extent, other basal forebrain pathways project via the fornix to entorhinal cortex. Basal forebrain projections via the temporal stem, amygdala, and entorhinal cortex to hippocampus are relatively unaffected by fornix transection (Gaffan, Parker, & Easton, 2001; Mesulam, Mufson, Levey, & Wainer, 1983; Ridley, Baker, Harder, & Pearson, 1996; Selden, Gitelman, Salamon-Murayama, Parrish, & Mesulam, 1998). Thus, AD's lesion interrupts major hippocampal output and the primary cholinergic input into the hippocampus, without affecting the structural integrity of the hippocampus and other MTL structures themselves. Although there are some efferent projections from the hippocampus to adjacent MTL structures via other pathways, AD's fornix lesion allows us to examine memory function while selectively removing a major contribution of the hippocampus and entorhinal cortex.

Bilateral damage to the fornix is known to impair anterograde memory, although there is considerable variability in the degree of memory loss that is reported (see reviews by Aggleton et al., 2000; Spiers, Maguire, & Burgess, 2001). Although some authors report that ablation of the anterior column of the fornix does not induce memory dysfunction (Bauer, Tobias, & Valenstein, 1993), others suggest that in certain cases, particularly after the removal of a colloid cyst, anterior fornicotomy may cause severe anterograde amnesia (Aggleton et al., 2000; Carmel, 1985; Easton, Ridley, Baker, & Gaffan, 2002; Gaffan, 1994; Gaffan, Gaffan, and Hodges, 1991; Garcia-Bengochea & Friedman, 1987; McMackin, Cockburn, Anslow, & Gaffan, 1995; Sweet, Talland, & Ervin, 1959), as it did in our patient. There are several reasons for such variability including, once again, the size and location of the lesion, type of memory tested, and whether the lesion is unilateral or bilateral (for review see Gaffan & Gaffan, 1991). There are far fewer studies on remote memory, but the available evidence suggests some loss, although the type and extent of remote memory loss is not well documented (D'Esposito, Verfaellie, Alexander, & Katz, 1995; Hodges & Carpenter, 1991; Park, Hahn, Kim, Na, & Huh, 2000; Spiers et al., 2001). Based on the purported effects of fornix lesions on anterograde memory, we predicted that remote episodic memory would be severely impaired. We also predicted that remote semantic would be relatively spared as has been reported in patients with fornix lesions (Spiers et al., 2001), and in patients with MTL/hippocampal lesions (for reviews see Moscovitch, Rosenbaum et al., 2005; Moscovitch,

Westmacott et al., 2005; Moscovitch, Nadel, Winocur, Gilboa, & Rosenbaum, 2006).

2. Case report

AD, a right-handed 45-year-old male, was referred to the Neurosurgery Department of a large Israeli medical center with a 1-month history of excruciating headaches. An MRI T1 study revealed a typical high-density non-enhancing colloid-cyst situated in the anterior part of the third ventricle, with a mild degree of obstructive hydrocephalus (see Fig. 1). The patient had no complaints of memory loss, and continued working up to the time of the surgery. Physical examination revealed no abnormalities.

Surgery was performed 1 month following his initial referral. A right frontal transcortical approach to the left Foramen of Monro, allowed the neurosurgeon access to the cyst. A typical dense colloid cyst was identified embedded in the anterior portion of the third ventricle, attached to its base. The cyst was removed via the Foramen of Monro with no unusual operative complications. AD's recovery in the immediate post-operative period was normal but soon after, he began to suffer from diabetes insipidus and became extremely confused.

Within 48 h of surgery, AD was interviewed by the first and second authors. He was unsure why he was hospitalized, could not recognize his neurosurgeon, and was unable to retain simple verbal material (such as three words) for more than a few minutes. AD's remote memory was also impaired. He was unable to recall any autobiographical events that occurred in the past 20 years. Although he recognized his wife and children, AD was unable to draw the exterior part of his house even though he had helped design it. His drawing of it was limited to a three-dimensional cube. After a 20-min bedside evaluation, the interviewers stepped out of the room. Upon their return, AD noted that he knew them, as he had met them "a few years ago". Two weeks following the operation, AD was re-evaluated and, although there was some general improvement, it was clear that he was severely amnesic.

2.1. Structural neuroimaging and lesion location

MRI brain scans were conducted 2 months post-surgery using an Elcint Prestige T2 machine with 5 mm thick scans. These scans showed no damage to the hippocampus and related medial temporal lobe (MTL) structures. An additional MRI was performed on the same machine, with 3 mm thick scans (TE 30/80, TR3000, 22 cm POV). An initial inspection of the resultant MRI scans documented a complete transection of the left anterior column of the fornix at the level of the anterior commissure, damage to the left anterior commissure, and a tiny infarct to the caudate nucleus on the right (Fig. 1). Later consultations with Dr. Fuqiang Gao from Sunnybrook Hospital in Toronto and Dr. Michael Alexander of Harvard Medical School confirmed these observations. Additional damage to the right anterior column of the fornix was identified which, given the limitations of the scans, was estimated to affect 75% of the tract, just anterior to the third ventricle above the anterior commissure. The

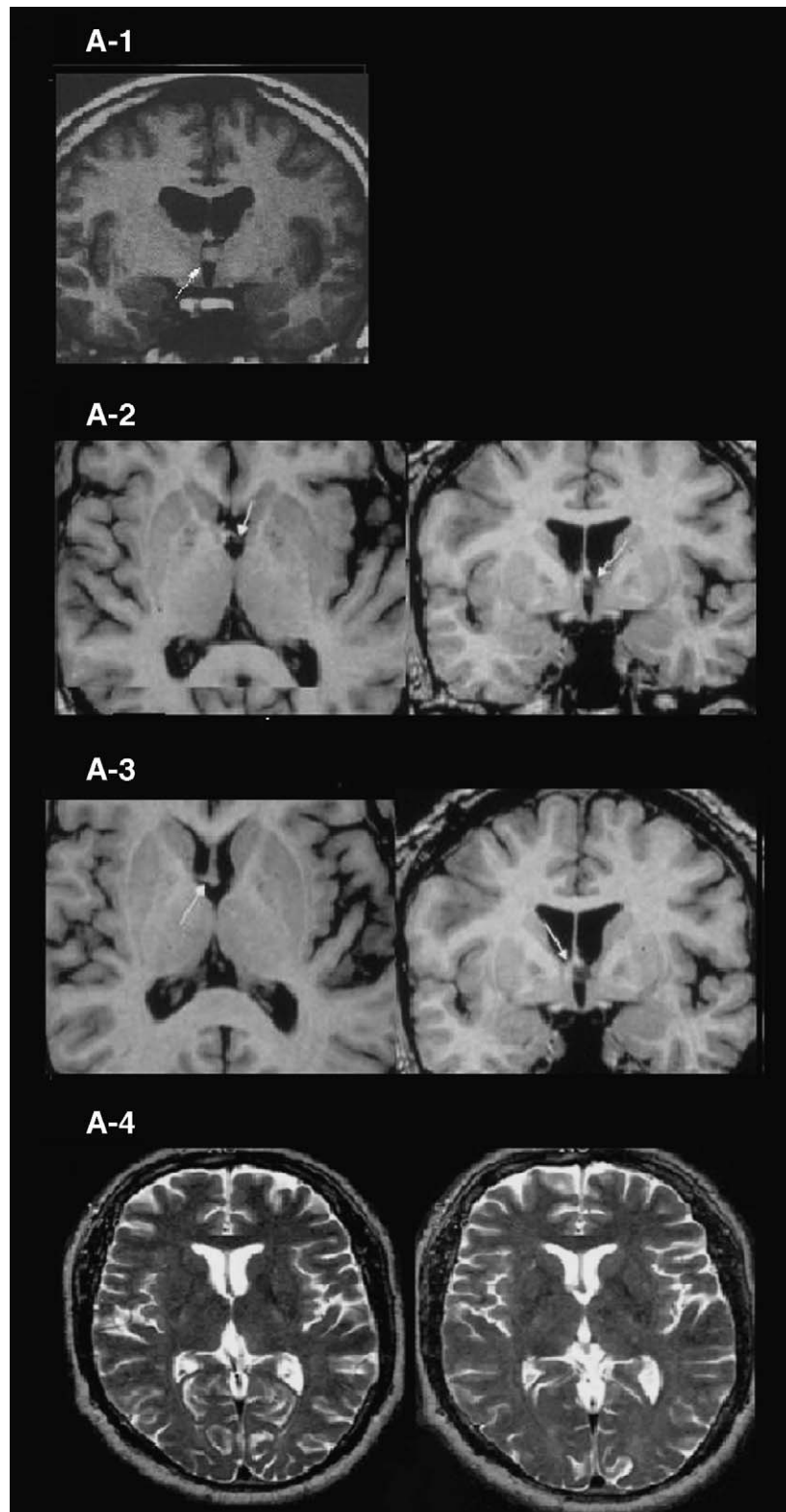


Fig. 1. Presurgical MRI scan of AD showing the colloid cyst (A-1). AD's postsurgical MRI and AD's bilateral fornix lesions can be seen in axial and coronal views of T1-weighted MRI's for the left fornix complete transection (A-2) and right fornix disruption of at least 75% (A-3). T2 weighted axial images further exemplify the fornix disruptions (A-4). AD's basal forebrain lesion can be seen in axial and coronal views (A-3) exemplifying the minimal extent of the lesion which involves midline basal forebrain nuclei on the left and possibly minimal right sided damage.

Table 1
Neuropsychological test results of AD 3 and 9 months postsurgery

Neuropsychological tests	Time of testing	
	3 months	9 months
Wechsler intelligence test III		(109) 55 percentile
Boston naming test	48 percentile	56 percentile
Wisconsin card sorting test		
Perseveration errors	6 percentile	1 percentile
Categories	5	4
Verbal fluency test		
Semantic	50 percentile	55 percentile
Phonetic	35 percentile	45 percentile
RAVLT learning curve	(6 6 7 7 6) 1 percentile	(4 7 5 8 9) 2 percentile
Postinterference	(1 + 2 intrusions) 1 percentile	(2 + 3 intrusions) 1 percentile
30 min delay	(9 + 8 intrusions) 1 percentile	(8 + 3 intrusions) 1 percentile
Spatial span		
Forward	(6) 50 percentile	(5) 50 percentile
Backward	(6) 50 percentile	(6) 50 percentile
Digit span		
Forward	(7) 50 percentile	(5) 50 percentile
Backward	(4) 50 percentile	(6) 50 percentile
Rey complex figure test		
Copy	(36/36) 85 percentile	(36/36) 85 percentile
Delay	(0/36) <1 percentile	(0/36) <1 percentile
Stroop test	6 percentile	–
Wechsler memory scale		
Logical prose immediate	(12.5/50) 1 percentile	(12/50) 1 percentile
Logical prose delay	(0/50) <1 percentile	(0/50) <1 percentile

medial part of the left basal forebrain showed a hypointense signal on the T1 and hyperintense signal on the T2, suggesting a very small lesion to the left medial basal forebrain, affecting the midline cholinergic nuclei of the basal forebrain, and sparing the lateral ones. SPECT brain scans using an Elcint XZ122 were also obtained. These scans were interpreted as showing a significant bilateral hypoperfusion of the mesial temporal regions.

2.2. Neuropsychological evaluation and anterograde memory

A month after the initial evaluation, AD became anxious and agitated about his condition. His wife noted that he often cried and wanted her to stay by his side. She also noted that each time he woke up, he reported that he was in a dream-like state. His MMPI-2 profile was elevated, with his scores on the Anxiety and Depression scales being $T=66$ and 67 , respectively. A formal neuropsychological evaluation was then conducted on an outpatient basis.

Neuropsychological test results are summarized in Table 1. The patient's intellectual performance, as measured by the WAIS-R, was in the higher end of the average range ($IQ=109$). AD's verbal abilities, as measured by the Boston naming test (BNT; Kaplan, Goodglass, & Weintraub, 1978), the controlled word association test (FAS) and semantic fluency (Benton, Hamsher, & Sivan 1983), were in the normal range. On the

Wisconsin card sorting test, another measure of executive functioning, AD performed within the normal range for his age, achieving five categories (WCST, Berg, 1948; Heaton, 1981) but he tested in the borderline range for number of perseverative errors. Informal testing did not reveal any dyslexic or dysgraphic errors. He had no difficulty in performing complex arithmetic tasks. Working memory, as measured by both digit and spatial span, was normal, as were visual constructional abilities, as measured by the Rey Osterrieth copy figure test (ROCF, Osterrieth, 1944).

In contrast to his relatively intact intellectual functioning, AD exhibited severe verbal and visual memory deficits. He showed no improvement with repeated presentation on the auditory verbal learning test (AVLT; Rey, 1958) and following interference, recalled only 1/6 of the words he had previously recalled. After a 30 min delay, he had no recollection of the task. On long-term recognition, his responses were at chance. On the Wechsler memory scale revised (WMS-R, Wechsler, 1981) story recall subtest, he scored in the impaired range even on immediate recall. After a 30 min delay, he was unable to recall having heard the story. When he was asked to recall the ROCF after a 3-min delay, AD's score was 0.

A follow-up neuropsychological evaluation was carried out 9 months post-surgery. During this testing, AD was also re-administered the MMPI-2 test. His profile indicated a heightened anxiety level and moderate depression. The results of the neuropsychological assessment are presented in Table 1.

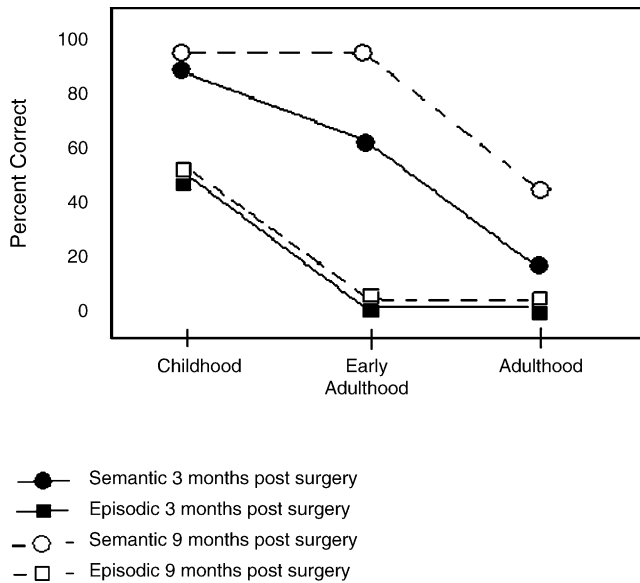


Fig. 2. Autobiographical memory interview (AMI) percent scores at childhood, early adulthood and adulthood. Percentages were calculated based on a maximum score of 21 for semantic memory, and 9, for episodic memory.

It can be seen that AD's anterograde amnesia persisted. His performance in other cognitive domains, including language, executive and working memory, also remained essentially unchanged.

2.3. Remote memory

AD's remote memory for public and autobiographical events was assessed to determine the nature and extent of his retrograde amnesia. Memory for faces and public events was assessed using the famous Israeli faces and famous Israeli public events tests developed by the first author. Items were drawn from commonly available books, scanned into PowerPoint presentation using a color scanner, and presented using a portable computer. Whenever AD could not answer a question, the examiner provided him with several multiple choice options.

Autobiographical memory was assessed by the Autobiographical Memory Interview (AMI, *Kopelman, Wilson, & Baddeley, 1989*) and a set of 16 family photographs that were provided by AD's daughter. All the stimuli were presented using a portable computer. The AMI consists of two sub-scales, Personal Semantic and Autobiographical Incidents, that sample memories from three time periods: childhood (ages 0–18), early adulthood (ages 18–30), and recent (within the past 5 years). The personal semantic sub-scale probes memory for general personal information such as names of friends and teachers, locations of schools attended, etc. The maximum score for each time period is 21 points. The Autobiographical Incidents sub-scale includes questions about specific events that require temporal and spatial contextual information for each incident. Three such incidents, such as "first day at work", are sampled at each time period with each incident receiving a maximum of 3 points, depending on the descriptive richness of the response and its specificity as to time and place.

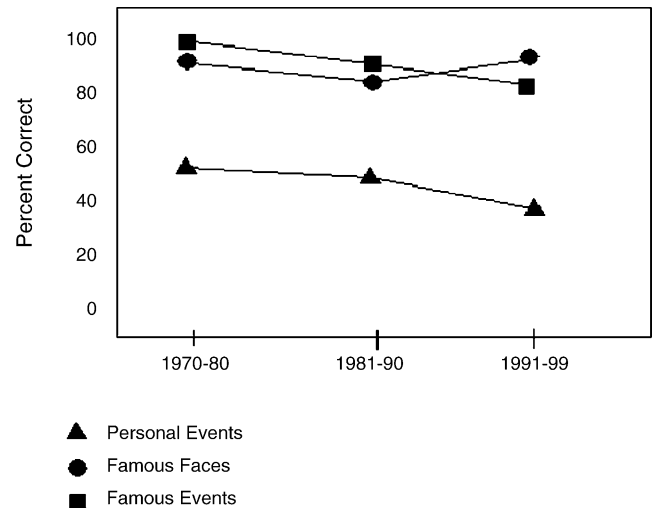


Fig. 3. Percentage correct for personal events (max = 16), famous faces (max = 19), and famous events (max = 20).

Fig. 2 depicts the results of the AMI. AD exhibited a general impairment in remote memory that was more severe for autobiographical incidents than for personal semantics. On the personal semantic sub-scale, AD showed a gradient in retrograde memory loss, in that the childhood score was within the normal range, whereas his early adulthood score indicated significant loss at 3 months which had recovered by 9 months. Scores for the recent time-period reflected an extensive anterograde memory deficit for personal semantics. Consequently, it is likely that memory for the most recent events draws on episodic memory as well as semantic knowledge. By contrast, memory for autobiographical episodes was severely impaired for the recent, and early adulthood periods, with borderline impairment of childhood memories. Interestingly, his childhood scores were elevated by two highly significant and emotional events that he recounts often and may constitute "personal folklore" (*Cermak, 1984; Cermak & O'Connor, 1983*) rather than true episodic memory.

On tests of famous faces and public events, AD was asked to name the person in the picture or identify the event during which the picture was taken. Whenever AD could not answer the question, he was prompted by a series of multiple choice options. For faces, there were 6, 6, and 7 faces from the 1970s, 1980s, and 1990s, respectively. For events, the corresponding numbers were 7, 6 and 7. He scored within the normal range for all time periods dating from the 1970s to the present. Performance on these tests was compared to that on a test of autobiographical memory in which he had to describe episodes depicted in 16 family photographs, 4, 6 and 6, for each decade. For each photograph, he was asked to provide details as to when the event occurred and who participated in the event. As seen in Fig. 3, he performed poorly on these tests relative to his performance on the famous faces and public events tests. We note, however, that he provided general semantic information related to the pictures (see below) without remembering the episodes to which they referred. His wife performed well on all of the measures and correctly identified more than 90% of the pictures. Below is a summary of some of his responses:

Photograph of an Army camp from the early 1970s

“Army base, these are T-type tanks.”

When was the picture taken? Why was it taken?

“I don’t know.”

Group photograph of a graduation ceremony in the early 1980s

“This is the graduation ceremony of some kind of training. I can’t remember the ceremony.”

Photograph of the circumcision of his son in the late 1980s

“This is a circumcision rite. I can see the baby.” (He didn’t identify his son).

A picture from a trip to Greece in the late 1990s

“A trip to Turkey.” (Wife said no).

“Rhodes?” (Wife said no).

“... we went on a train” (Wife said by boat).

3. Discussion

This study shows that bilateral lesions of the fornix lead to anterograde and retrograde amnesia without significantly affecting most other cognitive functions. The anterograde amnesia encompassed both verbal and non-verbal long-term memory in tests of recall and recognition, but short-term memory was spared. AD’s retrograde amnesia was more severe for autobiographical episodes than for personal semantics, thereby confirming the selective contribution of the hippocampal system and projections to basal forebrain to recovering remote episodic memory. There was a temporal gradient for personal semantics, with memories from early adult life and childhood being relatively spared. In contrast, memory for autobiographical episodes, even those from early childhood, was impaired. The latter results provide important new evidence for the dissociation between remote episodic and semantic memory with respect to hippocampal system function.

The impaired anterograde memory loss in AD resembles that observed following bilateral damage to the MTL, which includes the hippocampus (Eichenbaum & Cohen, 2001; Squire, Stark, & Clark, 2004) and its projections to basal forebrain (Gaffan et al., 2001). In most cases such damage is quite extensive and rarely restricted to the hippocampal formation. In our case, we demonstrated the typical features of MTL anterograde amnesia in a patient with lesions that were circumscribed to the fornical columns, and a small region of the basal forebrain, but left the rest of the MTL system, including the hippocampus, structurally intact.

Our findings are generally consistent with reports of anterograde memory loss following lesions of the anterior fornix, including those resulting from colloid cyst removal (Bauer et al., 1993; Calabrese, Markowitsch, Harders, Scholz, & Gehlen, 1995; Carmel, 1985; Gaffan & Gaffan, 1991; Gaffan et al., 1991; Garcia-Bengochea & Friedman, 1987; Spiers et al., 2001; Sweet et al., 1959; Tucker, Roeltgen, Tully, Hartmann & Boxell, 1988).

AD showed poor recall and poor recognition on the RAVLT, primarily because of an elevated false alarm rate. The impaired recognition differs from Aggleton et al’s (2000) report that patients with bilateral lesions to the fornix were only impaired on recall. Studies of animals with surgical lesions to the fornix have revealed the same pattern of preserved item recognition (e.g. Aggleton & Brown, 1999; Gaffan, Shields, & Harrison, 1984; Zola-Morgan, 1989) but impaired recognition of the same items when they were embedded in unique contexts (Gaffan, 1994). The deficit was increased when the basal forebrain was implicated (Gaffan et al., 2001). The reason for the discrepancy in AD’s results is not clear but it may be that AD tried to rely on general contextual cues rather than familiarity to support recognition.

By extension, the same arguments can be made with respect to retrograde amnesia following MTL damage. Two general patterns of retrograde amnesia have been reported in such patients. In one, there is a temporal gradient with recent memories being more severely affected, and very remote memories being spared, regardless of whether the memory is semantic or episodic. In the other, there is temporally extensive and severe retrograde amnesia, often dating back to early childhood, for episodic memory but a milder and temporally graded retrograde amnesia for semantic memory which is based more on familiarity (Fujii, Moscovitch, & Nadel, 2000; Moscovitch, Rosenbaum et al., 2005; Moscovitch, Westmacott et al., 2005; Moscovitch et al., 2006). AD’s retrograde amnesia conforms more closely to the second of the two patterns.

On the AMI, AD showed a temporally graded retrograde amnesia for personal semantics with early childhood memories being spared, and a more severe and temporally extensive deficit for autobiographical episodes. He had no autobiographical memories dating as far back as early adulthood, and had only a few, impoverished memories for early childhood. As we noted, for the latter period, his scores were elevated by two highly significant and emotional events that had been well-rehearsed and effectively had come to exist as semantic memories (Cermak, 1984; Cermak & O’Connor, 1983).

The results from the famous faces, famous public events, and family photos tests support this interpretation. Unlike the AMI, in which personal semantics may be contaminated by autobiographical episodes, the first two of the above tests are less prone to that confound. We found that, on the public events and faces test, AD scored normally at all the remote time periods, even the most recent ones, further indicating that remote semantic memory is not dependent on the hippocampal system. These findings also suggest that the temporal gradient for semantic memory observed in other patients with MTL damage may arise either from damage to extra-hippocampal structures, or from contamination by attempts to use episodic information related to the more recent events.

By contrast, on the family photos test, AD had a retrograde amnesia that extended to early childhood with no temporal gradient. The absence of a gradient on this test is especially informative because the nature of the test diminishes the possibility that well-rehearsed memories would be produced. Unlike the AMI, the memories elicited by the photos were not self-selected. As

such, they had to be based on reinstatement of autobiographical experiences in their spatial and temporal contexts rather than on “family folklore” (Cermak, 1984; Cermak & O’Connor, 1983).

Bayley et al. (2005) proposed that temporally extensive retrograde amnesia in MTL patients occurs only when the damage extends to lateral neocortex. Our results do not support this interpretation. Despite having no neocortical damage and small lesions to the fornix and basal forebrain area, AD’s retrograde amnesia for autobiographical episodes is temporally extensive and severe. It follows that the MTL, and the hippocampus in particular, are crucial for recovering autobiographical episodes, no matter how remote, and may also contribute temporarily to recovery of semantic memory. While there have been reports of patients with spared autobiographical memories following lesions restricted to MTL (Bayley et al., 2005), other investigators have found the amnesia to be far more extensive (Cipolotti et al., 2001; Moscovitch et al., 2006). The reason for these discrepancies has yet to be determined.

In addition to his fornix lesions, AD also has a small basal forebrain lesion. It is unlikely that this lesion alone caused his severe amnesia for the following reasons: (1) unilateral discrete lesions to the basal forebrain tend to produce modality specific (i.e. visual or verbal) deficits (Abe, Inokawa, Kashiwagi, & Yanagihara, 1998; Goldenberg, Schuri, Gromminger, & Arnold, 1999; Morris, Bowers, Chatterjee, & Heilman, 1992), whereas AD had both visual and verbal anterograde memory deficits. (2) Retrograde memory loss is minimal or non-existent following discrete basal forebrain lesions (Abe et al., 1998; Goldenberg et al., 1999; Hashimoto, Tanaka, & Nakano, 2000; Morris et al., 1992). (3) Finally, the precommissural columns of the fornix are the primary cholinergic routes from the basal forebrain to the hippocampus, with less extensive projections to neighboring MTL cortices (Easton et al., 2002; Morris et al., 1992). Given AD’s bilateral fornix lesions, no additional effect of basal forebrain lesions on hippocampal function is likely. The results of the SPECT scan, which show hypo-perfusion only in the MTL, are broadly consistent with this interpretation. However, it should be noted, that the basal forebrain lesion may exacerbate the effects of the fornix lesion (Gaffan et al., 2001). In this case, the only contribution that the basal forebrain lesion may have made to AD’s performance is an increased susceptibility to false alarms in recognition, and intrusions in recall (Abe et al., 1998; Goldenberg et al., 1999; Hashimoto et al., 2000; Morris et al., 1992). The important point to emphasize here is that only remote memory for autobiographical episodes was severely affected, whereas semantic memory was relatively spared. Although MTL extra-hippocampal structures such as the perirhinal cortex also receive cholinergic input from the basal forebrain, these are restricted to allocortical rather than neocortical regions, and they receive additional cholinergic input from lateral basal forebrain structures which were intact in AD. The relatively spared semantic memory believed to be mediated by them suggests that their function is sufficiently preserved.

The present findings add to the growing body of evidence supporting specialization within the MTL, with the hippocampus being crucial for re-experiencing episodic events (recollection) and extra-hippocampal structures, such as the entorhi-

nal and perirhinal cortex, for familiarity (Aggleton & Brown, 1999; Eichenbaum, 2001). Our finding that autobiographical memories, which rely on recollection, are impaired, whereas familiarity-based semantic memories are spared, indicate that this view applies as much to remote memory as to anterograde memory. These findings, together with converging evidence from other studies on humans (e.g. see Gilboa, Winocur, Grady, Hevenor, & Moscovitch, 2004; Moscovitch, Rosenbaum et al., 2005; Moscovitch, Westmacott et al., 2005; Moscovitch et al., 2006) and animals (Frankland & Bontempi, 2005; Rosenbaum et al., 2001; Winocur, Moscovitch, Fogel, Rosenbaum, & Sekeres, 2005), provide the basis for an emerging view, consistent with Multiple Trace Theory, that re-experiencing contextually rich memories are dependent on the hippocampal system no matter how old they are, whereas recovery of semantic memory can be accomplished independently of it (see Moscovitch, Rosenbaum et al., 2005; Moscovitch, Westmacott et al., 2005; Nadel & Moscovitch, 1997; Rosenbaum et al., 2001).

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References

- Abe, K., Inokawa, M., Kashiwagi, A., & Yanagihara, T. (1998). Amnesia after a discrete basal forebrain lesion. *Journal of Neurology, Neurosurgery & Psychiatry*, *65*, 126–130.
- Aggleton, J. P., & Brown, M. W. (1999). Episodic memory, amnesia and the hippocampal–anterior thalamic axis. *Behavioural and Brain Sciences*, *22*, 425–498.
- Aggleton, J. P., McMackin, D., Carpenter, K., Hornak, J., Kapur, N., Halpin, S., et al. (2000). Differential effects of colloid cysts in the third ventricle that spare or compromise the fornix. *Brain*, *123*, 800–815.
- Bauer, R. M., Tobias, B., & Valenstein, E. (1993). Amnesic disorders. In K. M. Heilman & E. Valenstein (Eds.), *Clinical neuropsychology* (3rd ed., pp. 523–602). New York: Oxford University Press.
- Bayley, P. J., Gold, J. J., Hopkins, R. O., & Squire, L. R. (2005). The neuroanatomy of remote memory. *Neuron*, *46*, 799–810.
- Bayley, P. J., Hopkins, R. O., & Squire, L. R. (2003). Successful recollection of remote autobiographical memories by amnesic patients with medial temporal lobe lesions. *Neuron*, *38*, 135–144.
- Benton, A. L., Hamsher, K., & Sivan, A. B. (1983). *Multilingual aphasia examination*. Iowa City: AJA Associates.
- Berg, E. (1948). Simple objective technique for measuring flexibility in thinking. *Journal of General Psychology*, *39*, 15–22.
- Calabrese, P., Markowitsch, H. J., Harders, A. G., Scholz, M., & Gehlen, W. (1995). Fornix damage and memory. A case report. *Cortex*, *31*, 555–564.
- Carmel, P. W. (1985). Tumours of the third ventricle. *Acta Neurochirurgica*, *75*, 136–146.
- Cermak, L. S. (1984). The episodic-semantic distinction in amnesia. In L. R. Squire & N. Butters (Eds.), *The neuropsychology of memory* (pp. 55–62). New York: Guildford.

- Cermak, L. S., & O'Connor, M. (1983). The anterograde and retrograde retrieval ability of a patient with amnesia due to encephalitis. *Neuropsychologia*, 21, 213–234.
- Cipolotti, L., Shallice, T., Chan, D., Fox, N., Scabhill, R., Harrison, G., et al. (2001). Long-term retrograde amnesia: The crucial role of the hippocampus. *Neuropsychologia*, 39, 151–172.
- D'Esposito, M., Verfaellie, M., Alexander, M. P., & Katz, D. I. (1995). Amnesia following traumatic bilateral fornix transection. *Neurology*, 45, 1546–1550.
- Easton, A., Ridley, R. M., Baker, H. F., & Gaffan, D. (2002). Unilateral lesions of the cholinergic basal forebrain and fornix in one hemisphere and inferior temporal cortex in the opposite hemisphere produce severe learning impairments in rhesus monkeys. *Cerebral Cortex*, 12, 729–736.
- Eichenbaum, H. (2001). The hippocampus and declarative memory: Cognitive mechanisms and neural codes. *Behavioural Brain Research*, 127, 199–207.
- Eichenbaum, H., & Cohen, N. J. (2001). *From conditioning to conscious recollection: Memory systems of the brain*. New York: Oxford University Press.
- Frankland, P. W., & Bontempi, B. (2005). The organization of recent and remote memories. *Nature Reviews Neuroscience*, 6, 119–130.
- Fujii, T., Moscovitch, M., & Nadel, L. (2000). Memory consolidation, retrograde amnesia, and the temporal lobe. In F. Boller & J. Grafman (Eds.) & L.S. Cermak (Section Ed.), *The handbook of neuropsychology: Vol. 2* (2nd ed.). Amsterdam: Elsevier.
- Gaffan, D. (1994). Dissociated effects of perirhinal cortex ablation, fornix transection and amygdectomy: Evidence for multiple memory systems in the primate temporal. *Experimental Brain Research*, 99, 411–422.
- Gaffan, D., & Gaffan, E. A. (1991). Amnesia in man following transection of the fornix: A review. *Brain*, 114, 2611–2618.
- Gaffan, E. A., Gaffan, D., & Hodges, J. R. (1991). Amnesia following damage to the left fornix and to other sites: A comparative study. *Brain*, 114, 1297–1313.
- Gaffan, D., Parker, A., & Easton, A. (2001). Dense amnesia in the monkey after transection of fornix, amygdala and anterior temporal stem. *Neuropsychologia*, 39, 51–70.
- Gaffan, D., Shields, C., & Harrison, S. (1984). Delayed matching by fornix-transected monkeys: The sample, the push and the bait. *Quarterly Journal of Experimental Psychology B*, 36, 305–317.
- Garcia-Bengochea, F., & Friedman, W. A. (1987). Persistent memory loss following section of the anterior fornix in humans. *Surgical Neurology*, 27, 361–364.
- Gilboa, A., Winocur, G., Grady, C. L., Hevenor, S. J., & Moscovitch, M. (2004). Remembering our past: Functional neuroanatomy of recollection of recent and very remote personal events. *Cerebral Cortex*, 14, 1214–1225.
- Goldenberg, G., Schuri, U., Gromminger, O., & Arnold, U. (1999). Basal forebrain amnesia: Does the nucleus accumbens contribute to human memory? *Journal of Neurology, Neurosurgery & Psychiatry*, 67, 163–168.
- Hashimoto, R., Tanaka, Y., & Nakano, I. (2000). Amnesic confabulatory syndrome after focal basal forebrain damage. *Neurology*, 54, 978–980.
- Heaton, R. K. (1981). *Wisconsin card sorting test manual*. Odessa, FL: Psychological Assessment Resources, Inc.
- Hodges, J. R., & Carpenter, K. (1991). Anterograde amnesia with fornix damage following removal of IIIrd ventricle colloid cyst. *Journal of Neurology, Neurosurgery & Psychiatry*, 54, 633–638.
- Kaplan, E., Goodglass, H., & Weintraub, S. (1978). *The Boston naming test. Experimental edition*. Philadelphia: Lea & Febiger.
- Kopelman, M. D., Wilson, B. A., & Baddeley, A. D. (1989). *The autobiographical memory interview*. Bury St. Edmunds: Thames Valley Test Company.
- McMackin, D., Cockburn, J., Anslow, P., & Gaffan, D. (1995). Correlation of fornix damage with memory impairment in six cases of colloid cyst removal. *Acta Neurochirurgica*, 135, 12–18.
- Mesulam, M. M., Mufson, E. J., Levey, A. I., & Wainer, B. H. (1983). Cholinergic innervation of cortex by the basal forebrain: Cytochemistry and cortical connections of the septal area, diagonal band nuclei, nucleus basalis (substantia innominata), and hypothalamus in the rhesus monkey. *Journal of Comparative Neurology*, 214, 170–197.
- Morris, M. K., Bowers, D., Chatterjee, A., & Heilman, K. M. (1992). Amnesia following a discrete basal forebrain lesion. *Brain*, 115, 1827–1847.
- Moscovitch, M., Nadel, L., Winocur, G., Gilboa, A., & Rosenbaum, R. S. (2006). The cognitive neuroscience of remote episodic, semantic and spatial memory. *Current Opinion in Neurobiology*, 16, 179–190.
- Moscovitch, M., Rosenbaum, R. S., Gilboa, A., Addis, D. R., Westmacott, R., Grady, C. L., et al. (2005). Functional neuroanatomy of remote episodic, semantic and spatial memory: A unified account based on multiple trace theory. *Journal of Anatomy*, 207, 35–66.
- Moscovitch, M., Westmacott, R., Gilboa, A., Addis, D. R., Rosenbaum, R. S., Viskontas, I., et al. (2005). Hippocampal complex contribution to retention and retrieval of recent and remote episodic and semantic memories: Evidence from behavioral and neuroimaging studies of healthy and brain-damaged people. In B. Uttl, N. Ohta, & C. Macleod (Eds.), *Dynamic cognitive processes: The fifth Tsukuba international conference*. Tokyo: Springer-Verlag.
- Nadel, L., & Moscovitch, M. (1997). Memory consolidation, retrograde amnesia and the hippocampal complex. *Current Opinion in Neurobiology*, 7, 217–227.
- Osterrieth, P. A. (1944). Le test de copie d'une figure complexe: Contribution à l'étude de la perception et de la mémoire (Test of copying a complex figure: Contribution to the study of perception and memory). *Archives de Psychologie*, 30, 206–356.
- Park, S. A., Hahn, J. H., Kim, J. I., Na, D. L., & Huh, K. (2000). Memory deficits after bilateral anterior fornix infarction. *Neurology*, 54, 1379–1382.
- Rey, A. (1958). *L'examen clinique en psychologie*. Paris: Presse Universitaire de France.
- Ridley, R. M., Baker, H. F., Harder, J. A., & Pearson, C. (1996). Effects of lesions of different parts of the septo-hippocampal system in primates on learning and retention of information acquired before or after surgery. *Brain Research Bulletin*, 40, 21–32.
- Rosenbaum, R. S., Winocur, G., & Moscovitch, M. (2001). New views on old memories: Re-evaluating the role of the hippocampal complex. *Behavioural Brain Research*, 127, 183–197.
- Selden, N. R., Gitelman, D. R., Salamon-Murayama, N., Parrish, T. B., & Mesulam, M. M. (1998). Trajectories of cholinergic pathways within the cerebral hemispheres of the human brain. *Brain*, 121, 2249–2257.
- Spiers, H. J., Maguire, E. A., & Burgess, N. (2001). Hippocampal amnesia. *Neurocase*, 7, 357–382.
- Squire, L. R., Stark, C. E., & Clark, R. E. (2004). The medial temporal lobe. *Annual Review of Neuroscience*, 27, 279–306.
- Sweet, W. H., Talland, G. A., & Ervin, F. R. (1959). Loss of recent memory following section of fornix. *Transactions of the American Neurological Association*, 84, 76–82.
- Tucker, D. M., Roeltgen, D. P., Tully, R., Hartmann, J., & Boxell, C. (1988). Memory dysfunction following unilateral transection of the fornix: A hippocampal disconnection syndrome. *Cortex*, 24, 465–467.
- Wechsler, D. A. (1981). *Wechsler adult intelligence scale-revised*. New York: The Psychological Corporation.
- Winocur, G., Moscovitch, M., Fogel, S., Rosenbaum, R. S., & Sekeres, M. (2005). Preserved spatial memory after hippocampal lesions: Effects of extensive experience in a complex environment. *Nature Neuroscience*, 8, 273–275.
- Zola-Morgan, S. (1989). Lesions of perirhinal and parahippocampal cortex that spare the amygdala and hippocampal formation produce severe memory impairment. *Journal of Neuroscience*, 9, 4355–4370.