Right temporofrontal cortex as critical locus for the ephory of old episodic memories

P Calabrese, H J Markowitsch, H F Durwen, H Widlitzek, M Haupts, B Holinka, W Gehlen

Abstract

A 54 year old patient of average intelligence with a severe and enduring loss of old autobiographical memories after herpes simplex type 1 infection is described. She was tested with a comprehensive neuropsychological battery two years after the infection. Special emphasis was laid on examining different aspects of retrograde memory. The neurological examination involved MRI and SPECT. Brain damage was found mainly in the right temporofrontal region, but minor left sided damage to this region seems possible. The patient was in the normal or slightly subnormal range for all tested anterograde memory functions, but manifested severe retrograde memory deficits with respect to episodic old memories and more moderate deficits in tests of general knowledge (semantic old memories). It is concluded that the ephory of old autobiographical memories relies heavily on an activation of the right lateral temporofrontal junction area, but that probably only some complementary left hemispheric damage to these regions will lead to major and persistent retrograde amnesia. Alternatively, the disconnection between major prefrontal and posterior cortical regions may provide a basis for retrograde amnesia.


Keywords: retrograde amnesia; memory; medial temporal lobe

In recent years evidence for an anatomical disassociation of anterograde and retrograde memory functions has accumulated. While the essential bottlenecks structures, relevant for information transmission for long term storage have largely been established, there is still a substantial lack of consistency with respect to the brain regions implicated in focal retrograde amnesia. The structures most often implicated in focal retrograde amnesia are situated in the prefrontal and temporal cortex, with some authors emphasising the importance of one structure or the other, and other authors assuming that only the combined and bilateral damage of portions of both frontal and temporal regions are necessary for the appearance of a fullblown retrograde amnestic syndrome. This inconsistency is enhanced by the currently widely accepted dissociation of different memory systems of which the episodic semantic distinction is the most common one. Episodic memories are those which can be traced back with respect to time and locus, whereas semantic memories signify those of a general nature (knowledge of the world).

In a recent review, the hypothesis that the combined action of inferolateral prefrontal and anteriorlateral temporal cortices is necessary for the retrieval of long term memory, was advanced on the basis of a comparison of several case reports and data on memory retrieval in normal subjects obtained with PET. Furthermore, it was speculated that these regions in the right hemisphere would primarily be engaged in triggering episodic old memories, whereas the same regions of the left hemisphere would be involved in retrieving information from the knowledge system. We report the case of a patient who fits into this proposed scheme by being selectively retrogradely amnesic in the episodic memory domain and having a SPECT documented hypoperfusion in the right temporofrontal junction area.

Case report

The patient, a 54 year old right handed woman had been admitted three years ago to the hospital as an emergency case with herpes simplex type 1 meningencephalitis.

NEUROLOGICAL AND NEURORADIOLOGICAL EVALUATION

The history of her illness started with complaints of nausea and tiredness one week before admission. Two days before admission she developed high fever with temperatures up to 39.8°C, complained about increasing headaches, and became more and more disoriented. Intermittently she was unable to recognise her family and showed signs of visual hallucinations for short periods. Finally she drifted into a state of severe drowsiness.

Her previous medical history involved left sided mastectomy for breast cancer as well as a hysterectomy, both about 10 years earlier. The family history of the patient disclosed no events with neurological, psychiatric, or other relevance.

On admission she was in a severe drowsy state, but could be awakened by painful external stimuli. She was then able to answer simple questions with yes or no, and to carry out simple movements to verbal command. A more complex form of communication, however, was not possible. Further neurological examination disclosed soft meningeal signs as well as a discrete to moderate left sided hemiparesis including
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Figure 1 General T1 weighted MRI after Gd DTPA contrast injection performed about three years after infection. Coronal sections from top left to bottom right demonstrate the principal loci of brain damage in the right inferior lateral prefrontal cortex (top left), the right temporopolar cortex (top right), and the right anterior temporal lobe (bottom left and right).

brachiofacial hemiparesis. Babinski’s sign was positive and the muscular reflexes were raised on the left. Reactions to painful stimuli could be elicited from both sides to the same extent.

An initial cranial CT showed a hypoattenuation in both hemispheres with a discrete right sided accentuation, predominantly in the frontotemporal region. After injection of an intravenous contrast a diffuse inhomogeneous enhancement was seen in the areas affected. A cranial CT control, one month later, was normal.

The first MRI was performed somewhat less than three months after onset of the disease. There was low signal intensity on T1 weighted images and high signal intensity on T2 weighted images in the right frontotemporal region with the maximum in the right temporal lobe, and, in addition, there was evidence of a circumscribed cystic substance loss in the right temporal and frontobasal brain regions. The damage included anterior portions of the right amygdalohippocampal region and affected in particular the region of the pathway of the right ventral branch of the uncinate fascicle (fig 1 in Markowitsch19). Results from a second MRI, performed about three years later, confirm those of the first (figs 1 and 2). Minor left sided

Figure 2 Axial proton density images through the centre of the patient’s brain damage, taken about three years after infection. Note that the extent of the brain damage closely follows the course of the ventral branch of the uncinate fascicle.
frontotemporal brain damage seems possible (figs 1 and 2).

An ECD SPECT, performed in the initial phase of the disease, showed a pronounced hyperperfusion in the right frontotemporal area. After three months, a control SPECT still showed a moderate hyperperfusion in the same region, whereas a control, done three years later, showed an area of circumscribed hypoperfusion in the same location (fig 3).

Initial EEG recording showed a moderate but generalised slowing of the background activity as well as a theta/delta focus in the right temporoparietal region. Furthermore, intermittent epileptogenic activity characterised by single sharp waves, was recorded in the same area, without clinical evidence of epileptic seizures during her stay in hospital. After six months the EEG recording was completely normal.

Analysis of CSF showed a white cell count of 840/3 cells and a protein level of 58 mg%. The serological data showed positive findings of IgG and IgM antibodies to the herpes simplex type 1 virus. Values for CSF became normal six months after onset of the disease.

NEUROPSYCHOLOGICAL EVALUATION
Testing was performed two years after the infection. The neuropsychological tests and a summary of the results are listed in the table. They included tests of intelligence, attention, concentration, language, cognitive flexibility, and various forms of memory tests. Most of these tests have been used and described in detail in previous publications.1 3 15 16

The patient was alert and motivated. She took part in conversations, was socially well adjusted, and was able to interact in various ways. When instructed, she could remember to do something or to go to a certain place. In formal testing, she gained an IQ of 103 points, and her attention was in the normal range. In the Wechsler memory scale R she was in the lower average range (general memory index 93 points). Other tests of verbal and non-verbal anterograde memory confirmed the impression of her normal to somewhat sub-average anterograde memory abilities. Her cognitive flexibility was below average.

RETROGRADE MEMORY
The patient described herself as lacking episodes of her personal past from several decades before her brain infection. We used the following tests to determine her performance level in the domains of retrograde episodic and semantic memory: the autobiographical memory interview (AMI)17 and the Crovitz form test1 were given to assess her
Neuropsychological tests used and main test results

<table>
<thead>
<tr>
<th>Assessed function and test</th>
<th>Patient</th>
<th>Normative data</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Control subjects</td>
</tr>
<tr>
<td>Intelligence</td>
<td>103</td>
<td>100 (SD 15)</td>
</tr>
<tr>
<td>Attention:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>d2 Concentration-endurance test</td>
<td>32%ile</td>
<td>25%ile-75%ile</td>
</tr>
<tr>
<td>Trail making test, form A</td>
<td>50%ile</td>
<td>50%ile</td>
</tr>
<tr>
<td>Language:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Token test</td>
<td>95%ile</td>
<td></td>
</tr>
<tr>
<td>Cognitive flexibility:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>FAS verbal fluency test</td>
<td>16</td>
<td>(cut off 17)</td>
</tr>
<tr>
<td>Trail making test, form B</td>
<td>83%  (25%ile)</td>
<td>50%ile</td>
</tr>
<tr>
<td>Proactive interference:</td>
<td>7, 6, 4, 4, 4*</td>
<td></td>
</tr>
</tbody>
</table>

Learning and anterograde memory:
- Digit span forward: 6, 4, 4, 5
- Digit span backward: 4, 4, 5
- Corsi block span forward: 5, 5
- Corsi block span backward: 4, 4, 5
- Wechsler memory scale-R:
  - Attention/concentration index: 87, 100
  - Verbal memory index: 99, 100
  - Visual memory index: 88, 100
  - General memory index: 93, 100
  - Delayed recall: 86, 100
- Rey auditory verbal learning test:
  - Trial 1: 6, 6, 4
  - Trial 2: 7, 8, 2
  - Trial 3: 9, 10, 2
  - Trial 4: 9, 11, 1
  - Trial 5: 11, 11, 6
  - Distractor (list B): 4, 4, 6
  - List 6: 100
- Rey complex figure:
  - Copy: 35, 35
  - 30 minute delay: 18, 18, 8
- Retrograde memory:
  - Autobiographical memory interview:
    - Personal semantic events (childhood, adulthood, recent): 14, 12, 5, 18, 21, 20, 21
    - Autobiographical incidents (childhood, adulthood, recent): 1, 2, 7, 8, 9, 5
  - Crovitz test (10 words per time period): See fig. 4
  - Famous faces test: 27%, 42%
  - Semantic general knowledge test: 71, 92

For the Rey auditory verbal learning test and the Rey complex figure means are given after Spreen and Strauss; for the AMI the scores are given for a control subject of comparable age (taken from Spreen and Strauss).  
*Although the number of remembered items is not atypical, the release from proactive interference (4, her last score) is below average.  
†See fig. 5.

episodic retrograde memory, and for testing her general knowledge the semantic general knowledge test of Schmidtke and a famous faces test were used. The AMI consists of a semistructured interview covering all epochs of a subject’s life; it is subdivided into two parts covering the more general, semantic, and the personal events domains. In the Crovitz form test 10 high frequency nouns are administered and the patient is asked to describe personal experiences and unique episodes relating to each word. The instructions were:

Figure 4. Performance of the patient in the Crovitz test. Out of 10 possible responses per period, the patient was able to give only three or two for her earliest decades, and five for the period after the infection.

Figure 5. Famous faces test. 15 faces were presented in periods. Here, the patient was less impaired than in autobiographical memory, but nevertheless showed a time gradient which reflects the onset of the infection in 1993.

"Relate personally experienced life events from any time period evoked by each of the following words . . .", and “Now estimate the date of its occurrence.” The semantic general knowledge test uses 100 items of the kind “What is the name for the child of a cow?” or “What is the currency used in Switzerland?” It furthermore includes questions on geography, famous people, and animals, animal products, and attributes. The famous faces test contained per decade 15 portraits of famous people (sportspersons, politicians, actors, etc).

By contrast with her abilities to learn and remember new information, she had persistent and severe deficits in her old memory abilities. They were most prominent for her autobiographical past (fig 4), whereas her general knowledge, measured with a famous faces test (fig 5) and the world knowledge test (and also partly reflected by the semantic part of the autobiographical memory interview) was much less affected (table).

Discussion

Our patient is another in a series of several patients with a clear dissociation between anterograde and retrograde memory abilities. She also fits into a series of patients with combined damage of predominantly temporopolar and inferolateral prefrontal regions and focal retrograde amnesia. (Her brain damage included the right medial temporal lobe as well, but we assume from comparing her case with previous ones that the lateral temporal lobe regions are critically implicated in retrograde amnesia.) Furthermore, she corresponds to the proposed scheme of hemispheric specific retrograde memory disturbances; her brain damage is predominantly right hemispheric and her memory disturbances are largely in the episodic memory domain. She also fits into the scheme proposed by Markowitsch for the occurrence of retrograde amnesia by having bilateral (although mainly right sided) brain damage. The etiology of her brain damage is not unusual for patients with severe memory...
disturbances,\textsuperscript{20, 21} including focal retrograde amnesia.\textsuperscript{22}

This case therefore combines all features of anatomo-behavioural interrelations for retrograde episodic amnesia, as predicted by Markowitsch's hypothesis.\textsuperscript{17} In addition, the overlap between episodic and semantic disturbances of old memories is in line with the assumption that these two memory systems have to be viewed more as a continuum than as sharply divisible entities: semantic information is generalised, repeated episodic information\textsuperscript{13} and the generalised (episodic plus semantic) deficiency of brain damaged patients at the stage of memory encoding\textsuperscript{24} may hold for the level of information retrieval as well, although it may be more distinct in the non-damaged brain.\textsuperscript{13}

The episodic memory system is most likely more sensitive to brain damage than the knowledge system: episodic information is unique,\textsuperscript{25} whereas semantic memories may derive from episodic ones through generalisation and repetition.\textsuperscript{23} Furthermore, episodic memories for ecphory probably need a synchronised activation of affect coding structures of the limbic system (amygdala).\textsuperscript{26} (Ecphory denotes the process by which retrieval cues interact with stored information so that an image or a representation of the information in question appears.)

The temporopolar orbitofrontal junction area will not be the locus of the engrams, but only a necessary mediator.\textsuperscript{13} Although we favour this region as the principal locus for memory ecphory, alternatively the disconnection between major prefrontal and posterior cortical regions may provide a basis for retrograde amnesia. Retrograde amnesia is certainly a multifaceted phenomenon which can accompany various kinds of diseases and which can occur after damage to quite divergent brain loci.\textsuperscript{14, 27}

\textbf{ROLE OF THE FRONTAL CORTEX IN RETRIEVAL}

Results from studies in patients with selective prefrontal damage usually do not indicate significant retrieval deficits. Instead, only the active, effortful engagement in ecphorising information may be disturbed.\textsuperscript{28} Other features of mnemonic information processing which are altered after prefrontal damage include aspects of metamemory and memory for the temporal order of events.\textsuperscript{29} Furthermore, the prefrontal cortex is viewed as acting in monitoring or supervising environmental stimulation, or as applying strategies.\textsuperscript{29}

All of these functions have, however, a closer affinity to the dorsolateral than to the orbitofrontal aspects of the frontal lobes. Damage to this region is assumed to induce changes in personality and emotional behaviour.\textsuperscript{29, 30} Brazzelli \textit{et al}\textsuperscript{15} described a patient with bilateral frontal and some additional circular and temporal damage who had severe amnesia, most likely including a failure to retrieve old memories although formal testing of retrograde memory could not be accomplished due to her incapacity to provide any kind of account. Recent \textit{O PET studies provide further evidence for a prefrontal involvement in verbal and pictorial episodic memory retrieval.}\textsuperscript{33}

In conclusion, the prefrontal cortex contributes to memory retrieval (or memory ecphory) both by providing the impetus or trigger for an active search of the engrams (stored at other places and most likely in a network-like fashion)\textsuperscript{13} and by its capacity as a time sensitive organiser. As mentioned above, autobiographical memory is composed of personally relevant and temporal knowledge. The temporal structuring or ordering of information is apparently necessary for its successful ecphory and retrieval. Results from single cases with prefrontal damage strongly support this view.\textsuperscript{33, 34} The patient's subnormal performance in all three tests on cognitive flexibility (and therefore a frontal lobe function) confirm the view, expressed by Kopelman,\textsuperscript{35} that test results in this domain are better predictors for retrograde memory performance than anterograde memory tests.

\textbf{ROLE OF THE TEMPORAL CORTEX IN RETRIEVAL}

Penfield\textsuperscript{36} has elucidated many of the phenomena occurring during electrical stimulation of the temporal lobe. One of his findings was the appearance of what he called "psychical responses": reproductions of past personal experiences which he attributed to the (lateral) temporal cortex. Recently Fink \textit{et al}\textsuperscript{17} similarly found with PET blood flow measurements a strongly increased activation in the anterior temporal regions in response to ecphorising autobiographical material. In summary, both case descriptions of epileptic patients with temporal lobe involvement and functional imaging data in normal human subjects indicate a role of lateral temporal portions in memory retrieval. The anterolateral temporal cortex most likely provides the connection to the posterior cortical centres of integration and therefore to the major storage places of the engrams.\textsuperscript{37}

\textbf{EMOTION AND MEMORY}

Some studies point to emotional changes after brain damage.\textsuperscript{15, 38, 39} The recent speculation that the right hemisphere is principally engaged in episodic memory retrieval and the left in semantic memory retrieval,\textsuperscript{13} is in line with such a view as episodic memories are usually more emotional than semantic ones. Autobiographical information is probably represented in a more complex way than semantic information and needs an involvement from the left language representing hemisphere as well as from the right emotional one. The case of an adolescent patient with automatic but not consciously reflected memory retrieval demonstrates this point.\textsuperscript{30}

It is not unlikely that the ecphory of episodic, but not of semantic, memory involves non-cortical limbic structures as well as the cortical ones, which primarily represent stored information.\textsuperscript{13} The basolateral limbic circuit, composed of portions of the prefrontal cortex, the amygdala, and the medial thalamus\textsuperscript{38} constitutes a basis for a network impli-
cated in emotional memory processing.19 41 The region of the medial thalamus can further be regarded as a central gating station for consciousness34 and therefore for the active, reflective processing, and associating of old memories. The mediodorsal nucleus not only projects to the prefrontal cortex, but especially to those portions of the temporal cortex as well which are implicated in the ephory of old memories—namely, the temporal pole (area 38).37 On the corticocortical level, the temporopolar and the ventrolateral prefrontal areas are bidirectionally interconnected by the uncinate fascicle.13 Emotional flattening is a frequent concomitant in patients with retrograde amnesia, both of organic and psychogenic origin.13 44 Taking all this evidence together, it is likely that damage to the temporofrontal junction areas may result in a blockade of access to stored engrams, particularly to those which require an emotional flavour for ephory as well.25 26

WHY IS THERE SUCCESSFUL RETRIEVAL OF MEMORIES ACQUIRED AFTER BRAIN DAMAGE WITHOUT THE RIGHT TEMPOROFRONTAL JUNCTION AREA? Patients with selective temporofrontal damage are still able to acquire new information long term after the insult and subsequently recall it long term.38 This recall must therefore occur via routes different from the damaged temporofrontal areas. The plasticity of the nervous system may lead to rewiring or rerouting of information acquired after the brain damage. Or the existence of hierarchically ordered recall systems can be assumed, similar to the models proposed for memory encoding after damage of relevant bottleneck structures.46 With respect to the differential retrieval of information acquired after brain damage (but not before brain damage), the additional requirement is that the process of information consolidation implies the immediate establishment of a retrieval path. In the intact brain this retrieval path would principally and primarily involve the temporofrontal junction areas. In the brain with damage to this region an immediate retrieval path would be established to other, intact structures, lower in hierarchy, and less accurate and precise in retrieval.

The proposal made by several authors implicitly or explicitly that the storage of the memory content is composed according to landmarks, may have the effect that after brain damage events which occurred before that landmark cannot be recalled, whereas those stored thereafter can be.4 This view might also be in accordance with Wolpaw’s that brain damage may disrupt the association between memories due to the “missing link” (in the temporofrontal junction area) which is necessary for the organised triggering (in the frontal portion) and access (in the temporal portion) to the engrams.35

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NEUROLOGY IN LITERATURE

Doctors’ writing, and writings

The quality of doctors’ writing has long been the object of public humour. I suspect that doctors, just as novelists, can suffer from writer’s block although in the case of the medical profession more perhaps from lack of time rather than from a failure of inspiration. Which explanation applies to Ovid Vere, in Heart and science, is not entirely clear. Thomas Mann’s encyclopaedia is one I have tried to emulate but as yet with no success in finding a willing publisher!

Samuel Johnson, 1779, Lives of the English poets—Dryden
It is a rare kind of giblet porridge, made of the giblets of a couple of young geese, stodged full of meteors, orbs, spheres, track, hideous draughts, dark characters, white forms, and radiant lights, designed not only to please appetite, and indulge luxury; but it is also physical, being an approved medicine to purge cholre: for it is propounded by Morena, as a receipt to cure their fathers of their choleric humours: and were it written in characters as barbarous as might be very well pass for a doctor’s bill.

Gustave Flaubert, 1856–7, Madame Bovary
Emma looked at him and shrugged her shoulders. Why hadn’t she at any rate one of those silent, earnest husbands who work at their books all night—and end up, by the time that rheumatism sets in at sixty, wearing a string of decorations on their ill-fitting dress-coats!

She would have liked this name of Bovary, that was hers, to be famous, on view at the book-shops, always cropping up in the papers, known all over France. But Charles had no ambition.

Wilkie Collins, 1883, Heart and science
His restless hand unlocked a drawer, and took out a manuscript work on medicine of his own writing. “Surely,” he thought, “I may finish a chapter, before I go to sea tomorrow?”

Thomas Mann, 1924, The magic mountain
Famous European specialists, physicians, psychologists, and economists will share in the composition of this encyclopedia of suffering, and the general editorial bureau at Lugano will act as the reservoir to collect all the articles which shall flow into it . . . This great work will not neglect the belletrist in so far as he deals with human suffering: a volume is projected which shall contain a compilation and brief analysis of such masterpieces of the world’s literature as come into question depicting one or other kind of conflict—for the consolation and instruction of the suffering.

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