Progressive degeneration of the right temporal lobe studied with positron emission tomography

P J Tyrrell, E K Warrington, R S J Frackowiak, M N Rosser

Abstract

A 79 year old man with a twelve year progressive history of prosopagnosia and recent naming difficulty, in whom other intellectual skills were preserved, is described. Positron emission tomography (PET) revealed an area of right temporal lobe hypometabolism, with an additional area of less severe hypometabolism at the left temporal pole. This may represent an example of progressive focal cortical degeneration similar to that associated with primary progressive dysphasia, but affecting the right temporal lobe.

Progressive, focal cognitive disorders due to localised cortical degeneration, have been described presenting as slowly progressive dysphasia,1 and less commonly, as dyspraxia, agnosia,2 or visual failure.3 The underlying cause, and the course, of these focal degenerations is unclear. They have been reported as both the initial presentation of primary degenerative dementia, and as a selective disorder progressing over many years, without evidence of more generalised cognitive impairment. While one case of focal progressive dysphasia eventually leading to dementia was shown at necropsy to have Pick's disease,4 5 a similar patient, with a more rapid progression to dementia, had the neuropathological features of Alzheimer's disease.6 Two further cases, who had no evidence of generalised dementia, showed cortical spongiform degeneration in the absence of Pick bodies, plaques or tangles.7 Metabolic studies with PET have shown left temporoparietal hypometabolism in two cases of progressive dysphasia without dementia.8

Although focal degenerations may affect other areas of cerebral cortex, involvement of the left temporal lobe is most commonly reported. The reason for this predilection is unknown, and examples of focal right temporal degeneration have not hitherto been reported. One of a series of six focal progressive dysphasia patients7 had evidence of minor right hemisphere hypometabolism, affecting particularly the anterior temporal lobe, in addition to the prominent left hemisphere hypometabolism. This patient had the longest history and was the most severely affected, which suggests that the non-dominant temporal lobe may be affected at a late stage.

The following case report describes a patient with a twelve year history of progressive prosopagnosia who had developed a naming deficit at a late stage. The main area of hypometabolism was in the right temporal lobe, with a less severe area in the left temporal lobe. This may represent an example of focal right temporal lobe degeneration.

Case history

A 79 year old right handed retired pharmacist presented with a progressive history of difficulty in recognising faces. Twelve years before presentation, he first noted impairment of his ability to remember the faces of people well known to him, which had progressed to the extent that he could no longer recognise his immediate family. In addition, for two to three years before presentation, he had difficulty remembering the names of places, and more recently difficulty with the names of friends and siblings and some difficulty naming objects. However, he continued to travel long distances alone, and to play competitive chess.

One half-brother became demented before death in his late eighties; there was no other family history of dementia or neurological disease. The patient's father had had pernicious anaemia. In the past, the patient had suffered a right-sided Bell's palsy. He was on thyroxine replacement for hypothyroidism, and vitamin B12 for pernicious anaemia.

On examination, he appeared very fit. General neurological examination was normal, apart from facial asymmetry without weakness. Speech was fluent with normal articulation, prosody and intonation. There was no evidence of apraxia, or primitive reflexes. General examination was normal, with a blood pressure of 160/90 mm mercury.

Haematological and biochemical investigations were normal, apart from the presence of positive gastric antibodies, and thyroid microsomal antibodies were positive at a titre of 1:6,400. Free thyroxine was within normal limits. Cerebrospinal fluid examination was normal, and syphilis serology negative. CT head scan showed minor enlargement of both sylvian fissures, particularly on the right (fig 1).

Neuropsychological assessment

On the WAIS-R he obtained a verbal IQ of 117 (pro-rated from four subtests), and a performance IQ of 101 (pro-rated from three subtests). On the Coloured Progressive Matrices he obtained a score (26/36 correct) close to the 75th percentile.10 (His scores on the above two tests are age-corrected). On Raven's Advanced Progressive Matrices Set A

Medical Research Council Cyclotron Unit, Hammersmith Hospital, London P J Tyrrell**
Department of Neurology, St Mary's Hospital, London R S J Frackowiak†
National Hospital for Nervous Diseases,† London E K Warrington
Correspondence to: Dr Tyrrell, MRC Cyclotron Unit, Hammersmith Hospital, London W12 0HS, United Kingdom
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Figure 1. CT head scan of the patient, showing enlargement of both sylvian fissures, most marked on the right. The left hand side of the scan corresponds to the right hemisphere in this figure.

his score (5/12 correct) was close to the 50th percentile. He obtained a reading IQ equivalent of 105 on the National Adult Reading Test.11

His score on the verbal version of the Recognition Memory Test (RMT) was 40/50 which is close to the 25th percentile.12 On the British Picture Vocabulary Test of single word comprehension his score (127/150) was above the 90th percentile.13 He had somewhat more difficulty in object naming tests. On the Oldfield Picture Naming Test he obtained a score of 18 correct for the first 20 items.14 On a more stringent Graded Naming Test his score was below the 5th percentile (3/30 correct) which is definitely impaired even when his age is taken into account.15 However, his comprehension of this vocabulary appeared to be intact as he obtained a high score in selecting the correct name from a choice of three (26/30 correct).

In contrast to his fairly satisfactory performance on a wide range of intelligence tests, verbal comprehension and verbal memory, his performance on a number of tests of visuospatial and visual memory was weak. Thus he had considerable difficulty in identifying incomplete letters (14/20 correct), and he identified only 6/20 unusual view objects, scores which are below the 5th percentile. He scored below the 5th percentile on an Objects Silhouettes Identification Test (3/15 correct).16 His recognition memory for faces (33/50 correct) although above chance, was below the 5th percentile.12 These impairments could not be attributed to sensory inefficiency as performance on shape discrimination (20/20 correct) and shape detection (30/30 correct) was entirely normal.17 His score on the RMT and all other tests quoted above are given in percentiles for the oldest age group for which norms are available (vis 55–70 years).

Although his scores on a number of tests of visual perception were somewhat weak, his performance on tests of face perception was quite creditable. On a faces matching test which required him to judge whether two photographs were of the same person or not, his score (16/20) was just below the 25th percentile. However, he scored at a normal level on the long version of Benton’s test of face perception (raw score 43 corrected for age and education).18

His most striking deficit was in recognising familiar faces. In identifying famous faces in a set of 12, he identified only one person (Mrs Thatcher). On a test of face familiarity on which the task is to choose the most famous person from two nonentity distractor stimuli, he obtained an exceptionally poor score (6/30 correct). On a verbal version of this test he obtained 27/30 correct.19 These findings corroborate the difficulty that he has in everyday life in recognising his family and friends by sight. He was considered to have a profound prosopagnosia.

Methods
15-O steady state positron emission tomography (PET) was performed on the CTI/931/08/12 scanner (Knoxville, Tennessee, USA) at the Medical Research Council Cyclotron Unit, Hammersmith Hospital, to obtain regional values of CMRO2 (cerebral metabolic rate for oxygen). Ethical approval for these studies, and for normal studies on eight volunteers (aged 59–83 years; mean 69.3 years), was obtained from the Ethical Committees of St Mary’s, the National and Hammersmith Hospitals.

Approval to administer radiolabelled gases was obtained from the ARSAC (Administration of Radioactive Substances Advisory Committee of the United Kingdom). Written consent was obtained from all patients and normal volunteers, after a full explanation of the procedure.

All normal subjects had no abnormalities on physical examination, and a normal blood pressure. They scored at least 28/30 on the Mini-Mental State Examination.20 Subjects with a history of seizures, significant head injury, hypertension, or neurological or other systemic disease were excluded. Two of the oldest volunteers (aged 74 and 83) had had CT scans performed, as part of another research project, which were normal.

The technique of 15-O-steady state scanning with correction for cerebral blood volume has been described previously,21–23 and was used to obtain parametric images of cerebral metabolic rate for oxygen (CMRO2), cerebral blood flow (CBF), oxygen extraction ratio (OER) and cerebral blood volume (CBV).

Images were inspected to ascertain the presence and position of focal deficits. Details of the image analysis technique has been published elsewhere,24 but essentially consists of placing standard anatomical stereotactic coordinates based on the Talairach atlas24 onto the parametric images, to obtain CMRO2 values for anatomical regions. Areas of hypometabolism in the patient were clearly delineated on the PET scan (fig 2). The regions of interest covered almost the whole width of the cortical ribbon. Visual inspection indicated that the areas of maximal hypometabolism were restric-
ted to the temporal lobes. Regions of interest were therefore placed on the anterior, middle, and posterior frontal gyri, and the superior, middle and inferior temporal gyri in the right and left hemispheres. Each structure was sampled with six regions of interest, from six adjacent anatomical planes. The mean value for each structure is presented. In addition, cerebellar cortex was sampled from two lower planes, and anterior and posterior parietal cortex from two representative planes through superior cuts distant from the visible sites of hypometabolism. Occipital cortex was sampled in three areas from three planes. Measurements were made from the left and right hemispheres independently.

Results
Regional values of CMRO₂ showed significant reductions in the superior temporal gyrus on both sides, more marked on the right than on the left; right: 1.47 ml/min/dl (mean value for the eight normal volunteers 2.76 ml/min/dl, standard deviation 0.41); left: 1.81 ml/min/dl (mean value for the eight normal volunteers 2.74 ml/min/dl, standard deviation 0.40). Values for middle and inferior temporal gyri, frontal cortex, parietal cortex, occipital cortex, cerebellum, and thalamus, caudate and putamen were within the normal range bilaterally.

Figures 2 and 3 show the CMRO₂ images of the patient, and a normal for comparison.

Discussion
The main area of hypometabolism relates to the right anterior temporal lobe, specifically in the right superior temporal gyrus. Clinical abnormalities arising from lesions of the non-dominant temporal lobe are often less apparent than the language deficits which are associated with dominant temporal lobe lesions. Poor performance on tasks of visual memory, as opposed to verbal memory are usually found with non-dominant lesions as was found in this case. The most striking and earliest abnormality in our case, however, was a profound prosopagnosia. It was found that the patient’s performance on a test of face perception was relatively satisfactory, whereas he had exceptional difficulty on tests of face familiarity judgement. Following the model of Bruce and Young it is inferred that his prosopagnosia can be attributed to damage to face recognition units: the individual’s familiar face vocabulary. Although it has been argued that bilateral lesions are required for prosopagnosia, there are many reported cases in which the syndrome arises from apparently unilateral right hemisphere lesions. Regions near the junction of temporal, occipital and parietal lobes are usually implicated, that is, more posterior than the area of maximal hypometabolism observed in our patient. In addition, the poor performance on visual perceptual tasks suggests an additional parietal lobe deficit not readily apparent on the CMRO₂ scan. Although a lesion in this area is indeed frequently implicated in cases of prosopagnosia, our findings raise the possibility that a prosopagnosia in which recognition mechanisms of familiarity are involved, (perceptual analysis of faces being preserved), may arise from a more anterior lesion.

The area of hypometabolism in the left anterior temporal lobe, associated with the observed nominal dysphasia in the presence of normal comprehension, is entirely consistent with data that implicates the left temporal lobe in word retrieval skills. The involvement of the left superior temporal gyrus alone, with preservation of the rest of the left temporal lobe, is also consistent with the findings of a common area of hypometabolism in the left superior temporal gyrus in the six cases with focal progressive dysphasia characterised by a naming deficit reported previously. The least affected of these patients, with a pure naming deficit, had hypometabolism restricted to this area.

The anatomical symmetry of the hypometabolism is of interest, with the most profound abnormality in the superior temporal gyrus on the right, and the only area of abnormality on the left confined to the superior temporal gyrus. This could be due to transcallosal connections, providing a route for transfer of a pathogenic agent. Alternatively, it may reflect selective vulnerability of the anterior temporal cortex to an unknown pathogenic agent or process. The most advanced case of primary progressive dysphasia reported previously showed a small area of hypometabolism in the right temporal lobe, in addition to severe hypometabolism in the left hemisphere.

The underlying aetiology of patients with progressive focal disorders is not known, but the very long progressive history in a normotensive man, and the absence of vascular lesions
on CT scan, argues in favour of a degenerative process. While Alzheimer's disease may present as a progressive dysphasia or as cortical blindness, the disease progresses rapidly to a generalised dementia. In addition, the PET scan in our patient did not show the posterior bipartical and bitemporal hypometabolism associated with dementia of the Alzheimer type, and the EEG was normal, in contrast to the slow wave abnormality usually seen in Alzheimer's disease.

Other cortical degenerations that may present as focal progressive cognitive disorders include Pick's disease, and unilateral Creutzfeld-Jakob disease. The underlying pathology may be similar to that described in cases of focal progressive dysphasia by Kirshner et al., with focal spongiform degeneration. While a number of cases of left temporal lobe degeneration have been described, right temporal lobe degeneration has not previously been reported. This may be due to selective vulnerability of the left temporal lobe, or that language disturbance may be more clinically apparent. It is noteworthy that the symptom which brought this patient to medical attention was the relatively recent onset of difficulty with object naming.

This case underlines the clinical and anatomical heterogeneity of the focal progressive cognitive degenerations, which presumably reflects heterogeneity of the underlying pathological processes.
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