Lateral parietal contributions to memory impairment in posterior cortical atrophy

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**A R T I C L E I N F O**

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Episodic memory
Attention
Alzheimer's disease

**A B S T R A C T**

Objective: Posterior cortical atrophy (PCA) is a neurodegenerative syndrome characterised by progressive impairment in visuospatial and perceptual function. Recent findings show that memory functioning can also be compromised early in the course of disease. In this study, we investigated the neural basis of memory impairment in PCA, and hypothesised that correlations would be observed with parietal cortex rather than classic medial temporal memory structures.

Methods: Eighteen PCA patients, 15 typical Alzheimer's disease (tAD) patients and 21 healthy controls underwent memory testing with the Rey Auditory Verbal Learning Test (RAVLT) word list and MRI. Voxel-based morphometry (VBM) was used to identify regions in the parietal and medial temporal lobes that correlated with memory performance.

Results: Compared with controls, PCA patients were impaired at learning, immediate and delayed recall and recognition of the RAVLT. Learning rate and immediate recall was significantly better in PCA compared to tAD, whereas there was no difference in delayed recall. Recognition memory also was not statistically different between patient groups, but PCA patients made significantly more false positive errors than tAD patients. VBM analysis in the PCA patients revealed a significant correlation between total learning and grey matter density in the right supramarginal gyrus, right angular gyrus and left postcentral gyrus. The left post central gyrus also significantly correlated with immediate and delayed recall and with recognition memory. No correlations were detected in the medial temporal lobe.

Conclusions: The findings provide novel evidence that early verbal memory impairment is frequently observed in PCA, and is associated with damage to lateral parietal structures. The results have implications for the diagnosis and management of PCA.

**1. Introduction**

Posterior cortical atrophy (PCA) is characterised by progressive impairment of visuospatial and visuoperceptual function that is not attributable to ocular disease (McMonagle et al., 2006; Tang-Wai et al., 2004). Impaired object and space perception is prominent, often accompanied by other features of posterior cortical dysfunction. The most common underlying cause is Alzheimer's pathology (Renner et al., 2004), although in a minority of cases alternative aetiologies, including corticobasal degeneration, dementia with Lewy bodies and prion disease, are implicated (Crutch et al., 2017).

Much research on PCA has concentrated on defining the visuospatial deficits (Crutch et al., 2013). Diagnostic criteria emphasise that episodic memory is relatively spared in the early stages (McMonagle et al., 2006; Tang-Wai et al., 2004).
Demographic and clinical characteristics of control and patient groups. Standard deviation given in brackets. Total scores achievable on neuropsychological tests, where applicable, in brackets in right column. Values in bold indicate significant group differences.

<table>
<thead>
<tr>
<th>Demographics</th>
<th>HC1</th>
<th>HC2</th>
<th>PCA</th>
<th>tAD</th>
<th>HC1 × PCA</th>
<th>HC1 × tAD</th>
<th>PCA × tAD</th>
<th>HC1 × HC2</th>
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<tbody>
<tr>
<td>N</td>
<td>21</td>
<td>45</td>
<td>18</td>
<td>15</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Age (yrs)</td>
<td>63.4 (6.1)</td>
<td>64.4 (7.2)</td>
<td>64.9 (6.8)</td>
<td>68.6 (9.7)</td>
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<td>0.126</td>
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</tr>
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<td>Education (yrs)</td>
<td>14.4 (2.1)</td>
<td>14.2 (3.0)</td>
<td>13.6 (2.0)</td>
<td>12.6 (6.5)</td>
<td>0.821</td>
<td>0.413</td>
<td>0.824</td>
<td>0.751</td>
</tr>
<tr>
<td>Gender (m:f)</td>
<td>9.12 / 25:20</td>
<td>9.12 / 25:20</td>
<td>9.7 / 25:20</td>
<td>9.8 / 25:20</td>
<td>0.656</td>
<td>0.535</td>
<td>0.849</td>
<td>0.336</td>
</tr>
<tr>
<td>Symptom duration (yrs)</td>
<td>–</td>
<td>–</td>
<td>3.8 (1.9)</td>
<td>3.0 (0.8)</td>
<td>–</td>
<td>–</td>
<td>0.130</td>
<td>–</td>
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<tr>
<td>DASS1 depression scale (normal range 0–9)</td>
<td>4.0 (4.5)</td>
<td>4.0 (4.5)</td>
<td>5.6 (5.0)</td>
<td>4.0 (4.5)</td>
<td>–</td>
<td>–</td>
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<tr>
<td>DASS anxiety scale (normal range 0–7)</td>
<td>3.4 (3.1)</td>
<td>3.4 (3.1)</td>
<td>9.1 (2.5)</td>
<td>9.1 (2.5)</td>
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<tr>
<td>DASS stress scale (normal range 0–14)</td>
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<td>0.019</td>
<td>0.019</td>
<td>0.019</td>
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</table>

**Table 1**

<table>
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<tr>
<th>Visual spatial function</th>
<th>HC1</th>
<th>HC2</th>
<th>PCA</th>
<th>tAD</th>
<th>HC1 × PCA</th>
<th>HC1 × tAD</th>
<th>PCA × tAD</th>
<th>HC1 × HC2</th>
</tr>
</thead>
<tbody>
<tr>
<td>ACE III Total (100)</td>
<td>96.0 (4.3)</td>
<td>96.0 (4.3)</td>
<td>84.9 (9.7)</td>
<td>77.6 (13.2)</td>
<td>0.000</td>
<td>0.000</td>
<td>0.092</td>
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</tr>
<tr>
<td>ACE III Attention</td>
<td>17.1 (1.5)</td>
<td>16.9 (1.5)</td>
<td>12.6 (4.0)</td>
<td>13.0 (3.0)</td>
<td>0.000</td>
<td>0.000</td>
<td>0.621</td>
<td>–</td>
</tr>
<tr>
<td>ACE III Fluency</td>
<td>24.7 (2.7)</td>
<td>24.7 (2.7)</td>
<td>18.4 (4.0)</td>
<td>15.5 (3.0)</td>
<td>0.000</td>
<td>0.000</td>
<td>0.465</td>
<td>–</td>
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<tr>
<td>ACE III Language (Zamboni et al., 2013)</td>
<td>25.7 (0.96)</td>
<td>25.7 (0.96)</td>
<td>22.7 (1.8)</td>
<td>19.1 (2.0)</td>
<td>0.000</td>
<td>0.055</td>
<td>0.153</td>
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</tr>
<tr>
<td>ACE III Visuospatial function (Uncapher and Wagner, 2009)</td>
<td>15.7 (0.72)</td>
<td>15.7 (0.72)</td>
<td>13.3 (3.0)</td>
<td>12.6 (2.8)</td>
<td>0.000</td>
<td>0.000</td>
<td>0.000</td>
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</tr>
</tbody>
</table>

**Abbreviations:** ACE III Addenbrooke’s Cognitive Examination; DASS Depression anxiety stress scale; HC1 Healthy controls group 1 (controls for behavioural data); HC2 Healthy control group 2 (controls for imaging data); PCA Posterior cortical atrophy; tAD typical Alzheimer’s disease; VOSP Visual object and space perception; RAVLT Rey auditory verbal learning task.

1 Missing data: Data in PCA patients was missing for some tests due to the test being ended on the patients’ request: Reduced sample sizes were present for: Rey figure (n = 9); VOSP cube analysis (n = 10); VOSP dot counting (n = 18); VOSP position discrimination (n = 11); PPT (n = 17); FAS (n = 17); Category fluency (n = 17); DASS (n = 14).

2006; Migliaccio et al., 2009). However, there is accumulating evidence of memory dysfunction in PCA (McMongle et al., 2006; Charles and Hills, 2005), and our work has shown that encoding and retrieval of new verbal information is significantly impaired in PCA patients compared to controls at initial presentation (Ahmed et al., 2016). Recently published consensus criteria also report that some PCA patients commonly report prominent memory disturbances at clinical presentation (Crutch et al., 2017). Since memory impairment could be amongst the presenting features in PCA, there is a need for better understanding of the neurocognitive mechanisms underlying this impairment, mechanisms which have to date remained entirely unexplored. Compared with tAD, voxel-based morphometry (VBM) studies in PCA show relatively minimal involvement of classically implicated memory circuitry i.e. the medial temporal lobes (MTL) (Kas et al., 2011; Lehmann et al., 2011; Whitwell et al., 2007). Instead, PCA patients show a characteristic pattern of atrophy in parieto-occipital and temporo-occipital cortices compared with controls, with significant grey matter reductions predominantly in the right superior parietal lobe (Lehmann et al., 2011).

Brain perfusion studies show bilateral and symmetrical hyperperfusion of the posterior cortex, with the most marked decrease found in the inferior parietal cortex (Kas et al., 2011).

Memory is not a cognitive function typically credited to the lateral parietal regions. Classic discourses describing the function of the parietal lobe make little mention of memory processes (e.g., Critchley, 1953; Luria, 1976). Recent investigation, however, suggests that the lateral parietal cortex may play an integrative role in episodic memory that has been largely underappreciated. Episodic memory tasks consistently show greater lateral parietal activation for encoding and retrieval of items, although more commonly for the latter process (Davidson et al., 2008; Hutchinson et al., 2009; Uncapher and Wagner, 2009), and lesion studies of patients with focal parietal damage provide further supporting evidence (Berryhill et al., 2007; Simons et al., 2010). Accordingly, given the relative sparing of the MTL in PCA, we hypothesize that memory impairment may be subserved by regions of the parietal lobe, and in particular, regions of the lateral parietal lobe that appear to be most severely compromised. This is in direct contrast to neural correlates of word list learning in AD. Using the Rey auditory verbal learning task (RAVLT), a widely used verbal episodic list learning task to quantify memory impairment in AD and preclinical AD, studies have shown consistent atrophy of medial temporal (Balthazar et al., 2010; Wolk and Dickerson, 2011) and medial parietal regions (Brugnolo et al., 2014) across imaging modalities.

The aim of this study was to characterise the behavioural and neuroanatomical profile of memory impairment in PCA. Specifically,
we predicted that PCA patients would not differ from AD patient controls in free recall measures of new learning but that PCA patients would benefit more from assessment of memory using recognition memory test formats, showing normal performance, given relative sparing of the MTL. Anatomically, we hypothesised that impairment on word list learning would correlate with sites of early and typical dysfunction in the lateral parietal regions in PCA, rather than classic memory structures of the MTL.

2. Material and methods

2.1. Participants

18 PCA patients were recruited through the Oxford Cognitive Disorders Clinic, Oxford, UK. Diagnosis was established by a senior behavioural neurologist (CRB, ST or MH) and neuropsychologists (IB and SA). All patients fulfilled consensus criteria for PCA (Tang-Wai et al., 2004; Crutch et al., 2017), based upon clinical assessment, brain imaging and detailed neuropsychological assessment. Clinical magnetic resonance imaging (MRI) confirmed focal atrophy in the occipital and parietal lobes.

Three controls groups were used for comparison with the PCA patients. The first group (HC1) consisted of 21 healthy controls, and was used to analyse neuropsychological test performance in the PCA patients. These participants had no objective cognitive impairment (scored > 88 on the Addenbrooke's Cognitive Examination-Revised (Mioshi et al., 2006)), and no prior history of psychiatric illness, significant head injury, or cerebrovascular disease, and were not prescribed any medication known to affect cognition.

The second group (tAD patient controls) consisted of 15 tAD patients, recruited from the dementia research clinic in Norwich, UK, as a disease control group for neuropsychological data analysis. tAD controls fulfilled consensus criteria for Alzheimer's disease (McKhann et al., 1984; McKhann et al., 2011), based upon clinical assessment, detailed neuropsychological assessment, and structural brain imaging. tAD patients showed marked impairment in episodic memory, with relatively preserved behaviour and personality, and characteristic bilateral medial temporal, with other more general atrophy.

PCA patients, HC1 and tAD control groups were matched for age, years of education and gender distribution (Table 1). PCA and tAD controls were matched for symptom duration, i.e. time since the first symptom was noticed. PCA patients did not demonstrate evidence of depressed mood or elevated stress levels, but did show significantly raised anxiety levels compared to controls. However, the group average (3.4) was still well within the normal range (0–7) on the Depression, Anxiety Stress Scale (DASS).

A third control group (HC2) consisted of a further 32 healthy participants, and was used solely for analysis of the PCA neuroimaging data. These controls were recruited from the Oxford Project to Investigate Memory and Ageing and the Memory and Amnesic Project, University of Oxford, UK (cohorts described fully in (Loane et al., 2018; Zamboni et al., 2013)). PCA patients and HC2 controls were matched for age (HC2: mean = 69.7 years, SD = 7.4; PCA: mean = 63.8 years, SD = 6.9; p > .05), education (HC2: mean = 14.5 years, SD = 3.5; PCA: mean = 13.7 years, SD = 1.9; p > .05), and gender distribution (HC2: 19 males, 13 females; PCA: 6 males, 7 females). HC1 and HC2 were also matched on age, education and gender (p > .05).

The study was approved by the National Research Ethics Service South Central - Hampshire B and Oxford C. All participants provided written informed consent in accordance with the Declaration of Helsinki.

2.2. Background neuropsychological tests

Standardised neuropsychological tests were administered to evaluate patient and control participant function in four domains:

(i) Global cognition: Addenbrooke's Cognitive Examination-III (Hsieh et al., 2013).
(ii) Visuospatial function: Dot counting, position discrimination and cube analysis from the Visual Object and Space Perception (VOSP; (Warrington and James, 1991) and the Rey-Osterrieth Complex fig. (29).
(iii) Visual imagery: Spatial Relations Test (adapted from (Kosslyn, 1987; Policardi et al., 1996)), Letter Shape Test (Farah, 1985) and Tail Judgement Test (Farah, 1988).
(iv) Language: Oral Pyramids and Palm Trees (PPT; (Howard and Patterson, 1992)), category fluency (Morris et al., 1989) and FAS letter fluency (Benton and Hamsher, 1976).

2.3. Memory assessment

2.3.1. Subjective memory questionnaires

PCA patients’ perspectives of their memory function were assessed using the Everyday Memory Questionnaire (EMQ; (Sunderland et al., 1984)) and carer perspectives were assessed using the Cambridge Behavioural Inventory-Revised (CBI-R; (Wedderburn et al., 2008)). Both patients and carers were instructed to answer questions with regards to memory and thinking and not poor visual functioning. For example, on the EMQ the last question asks how frequently the individual fails to recognise a close friend or relative – the results (Table 2) show that the majority of patients (93%) do not experience this problem in terms of memory loss, whereas if answered in terms of visual and perceptual symptoms in PCA, a higher proportion of patients would be expected to endorse this symptom. In both questionnaires, Likert scales were collapsed to provide a rating for symptoms experienced at three frequencies: Never, Frequent and Daily. Specifically, the CBI-R scales was collapsed as follows: “Never” = Never category; “a few times per month” and “a few times per week” = Frequent category; “daily” and “constantly” = Daily category. The EMQ scales was collapsed as follows: “not at all” = Never category; “about once in the last 3 months”, “about once a month” and “about once a week” = Frequent category; “about once a day” and “more than once a day” = Daily category.

2.3.2. RAVLT

The RAVLT (Rey, 1964) was used to measure objective memory performance. This well-validated instrument has been extensively employed to evaluate verbal episodic memory performance in various dementia syndromes (e.g. AD: (Balthazar et al., 2010; Wolk and Dickerson, 2011; Brugnolo et al., 2014); FTD: (Hornberger et al., 2010); PCA: (Charles and Hills, 2005)). A list of 15 unrelated words is read out to the participant five times, each time followed by free recall, in order to allow encoding of the word list. This is followed by an interference list of 15 new words. The original word list is then recalled immediately and at a delayed interval of 30 min. Recognition memory is assessed by presenting the participants with 50 words, consisting of 15 target words (i.e. the original word list) and 35 new words. If participants recognise a learned word, they respond “yes” (correct hit) and “no” to reject a new word. If participants incorrectly respond “yes” to a new word, it is scored as a false positive. A recognition memory index was calculated (hits - false positives) (Lemos et al., 2015; Grober et al., 2008) to describe recognition ability taking into account response bias. Values closer to the maximum 15 denote better recognition memory.

2.4. Structural neuroimaging

Image acquisition was conducted on a Siemens 3 T Trio system using a 32-channel head coil at the University of Oxford Centre for Clinical Magnetic Resonance Research. High resolution, 3D T1-weighted images were acquired using a Magnetization Prepared Rapid Gradient Echo (MPRAGE) sequence (echo time = 4.7 ms, repetition time = 2040 ms, 8° flip angle, field of view = 192 mm, voxel size = 1 × 1 × 1 mm). Scans from 5 PCA patients were removed due to
artifacts as a result of patient motion in the scanner. Structural data were analysed with an optimised protocol of the FSL-Voxel Based Morphometry (VBM) processing stream (see (Douaud et al., 2007)). In brief, images were brain extracted using the brain extraction tool (BET) (Smith, 2002) prior to tissue segmentation using the FMRIB Automatic Segmentation Tool (FAST). Images were non-linearly registered to standard space and an average study-specific template was created. All images were then non-linearly aligned to the study-specific template and modulated for correction of local field expansion or contraction by dividing the Jacobian of the warp field. Modulated images were then smoothed with an isotropic kernel with a sigma of 3 mm. Finally, voxelwise statistics were employed using a general linear model with non-parametric permutation testing (Winkler et al., 2014) and 5000 permutations were employed using the TFCE approach in Randomise with 5000 permutations per contrast and significance defined as p < 0.05. Age and gender were included as covariates in all analyses.

2.5. Statistical analyses

Demographic, clinical, and cognitive characteristics were explored using one-way analysis of variance with Sidak post hoc tests or independent samples t-tests, and Chi-squared tests for gender differences. RAVLT metrics were explored using Kruskal-Wallis nonparametric tests, with Mann-Whitney tests for pairwise comparisons. Spearman’s rank correlation coefficient was used to explore relationships between RAVLT metrics and clinical variables. Two-tailed tests were conducted with alpha level set at 0.05, with Bonferroni correction applied for multiple comparisons.

3. Results

3.1. Background neuropsychological assessment

PCA patients were impaired on all visuospatial and visual imagery tests compared to HCl, in keeping with the clinical phenotype of this syndrome, with some decline in language also (Table 1). Both PCA and TAD patients were impaired on the ACE-III compared to HCl, but there was no significant difference between patient groups.

3.2. Memory assessment

3.2.1. Subjective memory questionnaires

The most common observations endorsed by carers (those observed
by > 75% of carers as occurring frequently or daily) referred to symptoms related to memory retrieval and spatial memory. The most common subjective experiences of memory loss (those endorsed by > 75% of PCA patients) also pertained to memory retrieval, as well as task monitoring and memory for everyday activities (Table 2).

3.2.2. RAVLT

Both PCA and tAD patients learnt significantly fewer words across trials compared to HCl, and there was no significant difference between patient groups (Table 1). Both patient groups were impaired compared to HCl on immediate recall, but PCA patients recalled significantly more than tAD patients. To further explore encoding, rate of learning was computed (calculated as Trial 5 recall - Trial 1 recall; (Glosser et al., 2002)). Although both PCA and tAD groups showed significantly reduced rate of learning compared to HCl (PCA: p = .022; tAD: p = .040), PCA patients showed significantly more rapid learning than tAD patients (p = .040) (Fig. 1).

At retrieval, both patient groups were impaired compared to HCl on delayed recall, and there was no difference between patient groups. Examining recognition, PCA patients were comparable to HCl in correct identification of previously learned words (recognition hits), while the tAD group was impaired compared to HCl and PCA patients. Conversely, PCA patients produced significantly more false positives compared to HCl, while the tAD group showed no difference compared to HCl, although there was no significant difference between the patient groups. The recognition memory index revealed that, once the number of false positives was accounted for, both PCA and tAD groups were impaired compared to HCl, and the performance of both patient groups was comparable.

3.3. Other clinical considerations

Mood, symptom duration and impaired visual imagery were explored for their potential influence on poor memory performance in PCA.

3.3.1. Mood

There were no significant correlations between DASS stress, anxiety or depression measures with any of the RAVLT metrics (all p values > .05).

3.3.2. Symptom duration

To explore whether the memory profile in PCA was being driven by patients with a longer symptom duration who may have accumulated more widespread cognitive symptoms, PCA patients were divided into two subgroups based on median length of symptom duration (median = 3.3 years), replicating the method used by Kas et al. (Kas et al., 2011): (i) a short symptom duration group defined as having less than or equal to three years symptom duration (n = 9); and (ii) a long symptom duration defined as having greater than three years symptom duration (n = 9). There were no significant differences between these groups on any of the RAVLT metrics (all p values > .05). Supplementary analysis using symptom duration as a continuous variable revealed no significant correlations with the RAVLT.

3.3.3. Visual imagery

Poor visual imagery in PCA patients may impact the encoding process, based on evidence that visual strategies and mental imagery are commonly used to encode words presented in the auditory modality (Foley et al., 2010). There were no significant correlations between visual imagery tasks and RAVLT metrics.

3.4. Structural neuroimaging

3.4.1. Whole brain analyses

VBM analysis revealed significant decreases in grey matter density in PCA patients relative to HCl (Fig. 2). PCA patients showed characteristic pronounced changes, largely bilateral, in the lateral occipital cortex, and in lateral and medial parietal regions, including the parietal lobule, angular gyrus, the supramarginal gyrus, precuneus and posterior cingulate. Medial temporal regions were largely spared except for grey matter density reduction in bilateral posterior parahippocampal gyrus and posterior hippocampus (Table 3).

3.4.2. Region of interest analyses

Analyses within the parietal lobe mask revealed a significant correlation between total learning across trials and grey matter density in the right supramarginal gyrus, right angular gyrus and left postcentral gyrus. The left postcentral gyrus also significantly correlated with

![Fig. 1. Rate of learning (Trial 5 - Trial 1) across groups. Error bars represent standard error of the mean. (*p < .05).](image1)

![Fig. 2. Voxel based morphometry maps displaying reduced grey matter density in HC2 > PCA. The comparison was significant at p < 0.001 FWE corrected using TFCE method. Colour bars indicate T-values of statistically significant voxels.](image2)
immediate and delayed recall, correct recognition (Fig. 3). No correlations were detected between grey matter density within the parietal mask and the recognition memory index. No correlations with RAVLT metrics were detected within the MTL mask.

4. Discussion

This study provides novel evidence that, contra to traditional understanding, PCA is associated with early verbal memory impairment. Moreover, the degree of memory impairment correlates with atrophy in the lateral parietal cortex rather than regions typically associated with memory impairment, such as the MTL in tAD.

Compared with healthy controls, PCA patients were impaired at learning, immediate recall and delayed recall of the RAVLT word list. Nevertheless, PCA patients performed better than tAD patients on measures of learning rate and immediate recall, although their delayed recall was not significantly different. This overall impairment of verbal recall in PCA is in line with the subjective complaints of patients, and with our previous, retrospective study (Ahmed et al., 2016). Importantly, the results could not be explained as a function of disease progression or mood disorder. However, whilst we expected to see a relative sparing of recognition memory in PCA, this was not observed. Although PCA patients scored as many ‘hits’ as healthy controls, this was due to a response bias indicated by their greater number of ‘false positive’ responses. The reason for this response bias is unclear. A recent study suggests disinhibition may play a role, i.e. patients are unable to constrain endorsement of recognition items (Flanagan et al., 2016), although on the CBI-R, disinhibition did not emerge as a prevalent feature in PCA. An alternative possibility is that PCA patients have reduced confidence in their memory, akin to that observed in patients with parietal lesions (Simons et al., 2010), leading to a liberal response bias. Further studies are needed to explain this observation.

Investigating the neural substrate of the verbal memory deficit in PCA, we found that atrophy in lateral parietal regions correlated with both encoding and retrieval. Specifically, grey matter density in the right supramarginal gyrus, right angular gyrus and left postcentral gyrus correlated with total learning across trials, and the left postcentral gyrus with immediate recall, delayed recall and correct recognition. No correlations were identified between memory performance and MTL atrophy. This is strikingly different from extensive previous research in tAD, where learning and retrieval impairments have consistently been attributed to pathology in the hippocampus and surrounding cortex (e.g. (Balthazar et al., 2010; Wolk and Dickerson, 2011; Brugnolo et al., 2014)).

Two possible explanations for the observed neuroanatomical profile may be considered. First, the supramarginal gyrus has been associated with the phonological loop (Baddeley, 1986), which supports auditory-verbal working memory (Buchsbaum and D’Esposito, 2009). Accordingly, supramarginal gyrus atrophy in PCA may disrupt working memory processes essential in a verbal memory task. However, such processes are likely to be most critical to successful encoding of verbal information and therefore to learning of the initial trials on the RAVLT. For example, in tAD patients, Wolk and Dickerson (Wolk and Dickerson,

<table>
<thead>
<tr>
<th>Contrast</th>
<th>Regions</th>
<th>Cluster size</th>
<th>Coordinates</th>
<th>T value</th>
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</thead>
<tbody>
<tr>
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<td>Left: paracingulate cortex, paracingulate gyrus</td>
<td>98,166</td>
<td>56, −30, −4</td>
<td>7.86</td>
</tr>
<tr>
<td></td>
<td>Right: middle frontal gyrus</td>
<td></td>
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<td></td>
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<tr>
<td></td>
<td>Bilateral: superior parietal lobule, precuneal gyrus, supramarginal gyrus, postcentral gyrus, posterior cingulate, angular gyrus, parietal operculum cortex, Heschl’s gyrus, insular cortex, thalamus, superior temporal gyrus, middle temporal gyrus, inferior temporal gyrus, posterior parahippocampal gyrus, posterior hippocampus, putamen, caudate, precuneus, lingual gyrus, occipital fusiform gyrus, lateral occipital cortex, temporal fusiform gyrus, brain stem</td>
<td>98,166</td>
<td>56, −30, −4</td>
<td>7.86</td>
</tr>
</tbody>
</table>

Abbreviations: PCA Posterior cortical atrophy; HC2 Healthy controls group 2 (controls for imaging analysis).

immediate and delayed recall, correct recognition (Fig. 3). No correlations were detected between grey matter density within the parietal mask and the recognition memory index. No correlations with RAVLT metrics were detected within the MTL mask.

Fig. 3. Voxel based morphometry maps displaying correlation between reduced grey matter density in A = right supramarginal gyrus, right angular gyrus and left postcentral gyrus with total learning across trials; B = left postcentral gyrus with immediate recall; C = left postcentral gyrus with delayed recall and D = left postcentral gyrus with correct recognition in the PCA patients. Findings were significant at p < 0.01 FWE corrected using TFCE method (p values by colour bar).
found that reduced cortical thickness in the supramarginal gyrus correlated with learning on the initial trial of the RAVLT, while MTL structures correlated with total learning and delayed recall. In contrast, our results demonstrate that the supramarginal gyrus was associated with total learning and delayed recall in PCA patients, suggesting that impaired auditory-working memory is unlikely to be the primary deficit underpinning the memory impairment.

A second and more compelling possibility is that the lateral parietal correlates of memory in PCA reflect the hitherto under-recognised significance of these regions in episodic memory. Classic theories of memory would hypothsize that, because MTL involvement is not an early feature of PCA, patients should have relatively intact long-term declarative memory. However, an increasing number of fMRI (e.g. (Cabeza and Nyberg, 2000) and lesion (e.g. (Davidson et al., 2008) studies have demonstrated a role for bilateral parietal cortex in both the encoding and retrieval of memories. The parietal cortex is also firmly linked with attentional processing and is the most anterior of the neurophysiological, neurophysiological, and neuroimaging literature that explores memory impairment in PCA. These systems were adminstered to the patient groups in visual imagery performance and its potential impact upon memory. Additional tests of attention and working memory could be more systematically studied using free and cued memory tests (e.g. (Teichmann et al., 2017)) that control encoding and retrieval of the material to be learnt.

5. Conclusions

In summary, the results of this study provide the first description of the neural correlates of memory impairment in PCA, and strongly suggest that the deficits are underpinned by damage within parietal rather than medial temporal lobe networks. Future work is warranted to explore the relationship between attention and memory in PCA.

Competing interests

None.

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