Accepted Manuscript

Evidence of an Amnesia-Like Cued-Recall Memory Impairment in Nondementing Idiopathic Parkinson's Disease

N.M.J. Edelstyn, C.M. John, T.A. Shepherd, J.L. Drakeford, D. Clark-Carter, S.J. Ellis, A.R. Mayes

Please cite this article as: Edelstyn NMJ, John CM, Shepherd TA, Drakeford JL, Clark-Carter D, Ellis SJ, Mayes AR, Evidence of an Amnesia-Like Cued-Recall Memory Impairment in Nondementing Idiopathic Parkinson's Disease, *CORTEX* (2015), doi: 10.1016/j.cortex.2015.06.021.

1

2

3 **Evidence of an Amnesia-Like Cued-Recall Memory Impairment**

4 **in Nondementing Idiopathic Parkinson's Disease.**

5

6 N.M.J. Edelstyn*[1], C.M. John[1], T.A. Shepherd[1],

7 J.L. Drakeford[2], D. Clark-Carter[2], S.J. Ellis[3] and A.R. Mayes[4].

8

9 [1]School of Psychology, Keele University, Staffordshire. UK.

10 [2]School of Psychology, Sport and Exercise, Staffordshire University, Staffordshire. UK.

11 [3]University Hospital of North Midlands and Keele University, Staffordshire. UK.

12 [4]Psychological Sciences, University of Manchester, Manchester. UK.

13

14 **Running title:** Cued-recall impairment in Parkinson's Disease.

15

16 ***Corresponding author:** Nicky Edelstyn, School of Psychology, Keele University, Keele,

17 Staffordshire ST5 5BG, UK. Tel.0044 1782 734318. Email: n.edelstyn@keele.ac.uk

18

1 **Abstract**

2 Medicated, non-dementing mild-to-moderate Parkinson's disease (PD) patients usually show

3 recall/recollection impairments but have only occasionally shown familiarity impairments. We aimed

4 to assess two explanations of this pattern of impairment. Recollection typically improves when

5 effortful planning of encoding and retrieval processing is engaged. This depends on prefrontally-

6 dependent executive processes, which are often disrupted in PD. Relative to an unguided encoding

7 and retrieval of words condition (C1), giving suitable guidance at encoding alone (C2) or at encoding

8 and retrieval (C3) should, if executive processes are disrupted, improve PD recollection more than

9 control recollection and perhaps raise it to normal levels. Familiarity, being a relatively automatic

10 kind of memory, whether impaired or intact, should be unaffected by guidance. According to the

11 second explanation, PD deficits are amnesia-like and caused by medial temporal lobe dysfunction

12 and although poorer recollection, which is caused by hippocampal disruption, may be improved by

13 guidance, it should not improve more than control recollection. Familiarity impairment will also

14 occur if the perirhinal cortex is disrupted, but will be unimproved by guidance. Without guidance,

15 recollection/recall was impaired in thirty PD patients relative to twenty-two healthy controls and

16 remained relatively equally impaired when full guidance was provided (C1 vs C3), both groups

17 improving to broadly the same extent. Although impaired, and markedly less so than recollection,

18 familiarity was not improved by guidance in both groups. The patients showed elevated rates of

19 subclinical depressive symptoms, which weakly correlated with recall/recollection in all three

20 conditions. PD executive function was also deficient and correlated with unguided/C1 recollection

21 only. Our results are consistent with a major cause of the patients' recall/recollection impairments

22 being hippocampal disruption, probably exacerbated by subclinical depressive symptoms. However,

23 the results do not exclude a lesser prefrontal cortex contribution because patient executive

24 functions were impaired and correlated solely with unguided overall recollection.

1   **Key words:** Recall/recollection, Familiarity, medial temporal lobe, prefrontal cortex,

2   Parkinson's Disease.

3

1 Introduction

2 Idiopathic nondementing PD is dominated by tremor, bradyphrenia, rigidity and postural instability

3 (Parkinson, 1817), which each have a good response to dopaminergic medication. The motor

4 problems that characterise PD are caused by progressive degenerative changes that occur primarily

5 in the dopaminergic nuclei of the midbrain (Hornykiewicz, 1966; Fearnley & Lees, 1991; Obeso, *et*

6 *al.,* 2008). However, PD is not 'just'a motor disorder. Patients are often impaired at recalling recently

7 experienced information, such as facts and personal events, but have usually been found to be less

8 impaired at recognition of recently encountered stimuli (e.g., Flowers, Pearce & Pearce, 1984;

9 Taylor, Saint-Cyr & Lang, 1986; Breen, 1993; Beatty, Staton, Weir, Monson & Whitaker, 2003;

10 Higginson, Wheelock, Carroll & Sigvardt, 2005; Edelstyn, Mayes, Condon, Tunnicliffe & Ellis, 2007;

11 Edelstyn, Shepherd, Mayes, Sherman & Ellis, 2010; Shepherd, Edelstyn, Mayes & Ellis, 2013).

12 The most plausible explanation of why recall is typically more impaired than recognition in PD is

13 provided by the widely accepted dual process view according to which recognition is supported by

14 two kinds of memory, recollection and familiarity, that depend on distinct processes and different

15 systems of brain structures (see Montaldi & Mayes, 2010; but also see Wixted & Squire, 2011 for a

16 contrasting view). Recollection is a kind of memory in which a recognition test stimulus cues recall of

17 detail(s) from one or more previous episodes where the stimulus was encountered. This kind of cued

18 recall is often diagnostic of an earlier encounter with the stimulus. It is functionally and neurally very

19 similar to other kinds of recall, which also depend on cues (even free recall is cued by context). Tests

20 of recall, therefore, depend solely on recollection or kinds of recall very similar to it whereas

21 recognition test performance can often be strongly supported by familiarity. Familiarity-driven

22 recognition memory supports a subjective experience of memory for a previously encountered

23 stimulus in the absence of any cued recall of associated details from previous encounters with the

24 stimulus. Whereas recollection and other kinds of recall often depend on effortful and planned

25 processing at study and test, familiarity is believed to be a relatively automatic activity (see Jacoby,

4

1   1991; Yonelinas, 2002). In general, the dual process view explains the typical PD memory

2   impairment as being the result of a brain dysfunction in a region that primarily mediates recollection

3   and other kinds of recall, leaving the familiarity brain system functioning relatively normally. The

4   dual process view can also explain cases where recognition is clearly impaired because, although

5   familiarity often makes a major contribution to recognition, the relative contribution of recollection

6   and familiarity to different kinds of recognition varies and some kinds mainly depend on recollection

7   (Holdstock, Mayes, Roberts, Cezayirli, Isaac & O'Reilly, 2002).

8   Two hypotheses are consistent with PD patients having a greater recall than recognition deficit. Both

9   hypotheses propose that the PD memory disorder is caused by dysfunction (but also possibly

10   damage) of brain regions that are modulated by the dopaminergic midbrain structures the atrophy

11   of which underlies PD. According to the prefrontal cortex/organizational hypothesis, the death of

12   dopamine-producing cells in the *ventral tier* of the substantia nigra pars compacta (Fearnley & Lees,

13   1991) results in a progressive loss of dopamine innervation in the basal ganglia (Hornykiewicz, 1966).

14   Since a close relationship exists between the basal ganglia and prefrontal areas (Alexander, Delong &

15   Strick, 1986), it is not surprising to find evidence of PD-dependent deficits of forms of cognition, such

16   as working memory (Gabrieli, Singh, Stebbins & Goetz, 1996), planning and problem-solving (Morris,

17   Downes, Sahakian, Evenden, Heald & Robbins, 1988; Beatty & Monson, 1990), and verbal fluency

18   (Hanes, Andrewes, Smith & Pantelis, 1996), that depend upon the integrity of the prefrontal cortex

19   (Owen, Doyon, Dagher, Sadikot & Evans, 1998). Common to each of these kinds of cognition is the

20   requirement to select and implement appropriate strategies. Evidence of poor performance of

21   these executive processes (for a review see Dirnberger & Jahanshahi, 203) and functional brain

22   imaging evidence of reduced (medial) prefrontal activation and elevated false alarm rate during a

23   yes/no item recognition memory task in PD (e.g., Segura, *et al.*, 2012) has led to proposals that the

24   prefrontal dependent executive deficits underlie the breakdown of recall, which often depends on

25   effortful and organized encoding and retrieval processes. The impaired executive control processes

26   normally underlie the organization of material at encoding and the search for better cues at retrieval

5

1 as well as retrieval-related activities, such as response monitoring and decision-making (e.g.,

2 Gabrieli, *et al.,* 1996; Mayes & Daum 1996; Savage, *et al.,* 2001).

3 The claim of the prefrontal cortex / organisational hypothesis that prefrontal dysfunction disrupt

4 recall is supported by findings from lesion patients. Like PD patients, patients with frontal lesions are

5 not amnesic; however, they do exhibit deficits on tests of free recall (Gershberg & Shimamura, 1995;

6 Wheeler, Stuss & Tulving, 1995; Turner, Cipolotti, Yousry & Shallice, 2007) and source recall,

7 particularly when the tests depend heavily on the use of memory strategies as applies when multiple

8 learning trials are used (Gershberg & Shimamura, 1995; Janowsky*,* Shimamura & Squire, 1989a;

9 Janowsky, Shimamura, Kritchevsky & Squire, 1989b; Duarte, Ranganath & Knight, 2005). On the

10 other hand, when the demands on effortful executive control processes are reduced by the

11 provision of category cues at test and / or strategic instruction, recall performance has been shown

12 to be less affected (Gershberg & Shimamura, 1995); and item recognition memory is relatively

13 normal (Janowsky *et al.,* 1989a; Shimamura, Jurica, Mangels & Gershberg, 1995; Parkin*,*

14 Bindschaedler, Harsent & Metzler, 1996; Schacter, Curran, Galluccio, Milberg & Bates, 1996). These

15 findings suggest that a deficit in the initiation and use of organisational strategies   explains the free

16 recall deficits following frontal lobe damage. If this is so, then provision of strategic instruction at

17 encoding and/or retrieval should remediate these deficiencies as has been reported (Jetter, Poser,

18 Freeman & Makowitsch, 1986; Hirst & Volpe, 1988; Della Rocchetta & Milner, 1993; Gershberg &

19 Shimamura, 1995).

20 Although prefrontal damage can disrupt familiarity (e.g., Duarte et al., 2005; MacPherson, Bozzali,

21 Cipolotti, Dolan, Rees & Shallice, 2008), and fMRI studies also indicate that activation of various

22 prefrontal regions occurs during familiarity processing (e.g. Henson, Rugg, Shallice, Josephs & Dolan,

23 1999), there is very poor correspondence of prefrontal sites between the two approaches and

24 relatedly what they are doing. There is also behavioural evidence from older healthy adults showing

25 better executive functioning is associated with greater recollection, but is unrelated to familiarity

6

1 performance (Anderson, Ebert, Jennings, Grady, Cabeza & Graham, 2008). This is consistent with the

2 view that familiarity is a much more automatic kind of memory and depends much less on

3 executive function than intentional recollection, and consequently, the prefrontal/ organizational

4 hypothesis would predict that familiarity should be relatively spared, as has been reported (e.g.,

5 Mayes & Daum, 1996).

6 The second hypothesis that explains why recall is typically more impaired than recognition in PD

7 proposes that hippocampal dysfunction selectively disrupts recall memory in the same way that it is

8 disrupted in organic amnesia. This hypothesis is derived from the neuroanatomical development of

9 the dual process view (e.g., Aggleton & Brown, 1999), a central tenet of which is that the

10 hippocampus and its connections via the fornix in the midline diencephalon and the retrosplenial

11 cortex play a selective role in mediating recollection and other kinds of recall. Based on this, the

12 medial temporal lobe / amnesia-like hypothesis proposes that impaired dopaminergic modulation

13 predominantly of the hippocampus is caused by atrophy of the *dorsal* tier of the substantia nigra

14 pars compacta and the ventral tegmental area (Fearnley & Lees, 1991; Lisman & Grace, 2005;

15 Bunzeck, *et al.,* 2007). The hypothesis also allows that direct neuropathology of the hippocampus

16 may occur in PD. This neuropathology may involve several hippocampal subregions and, particularly

17 if PD is accompanied by subclinical depressive symptoms or clinical depression, probably includes

18 suppression of neurogenesis in the dentate gyrus (for a full discussion of adult hippocampal

19 neurogenesis in depression, see Kempermann & Kroneberg, 2003; Sahay & Hen, 2007). However,

20 whether these neuropathologies result directly and solely from atrophy of midbrain dopaminergic

21 structures remains uncertain (e.g., Laakso, *et al.,* 1996). Whatever the source of dysfunctionality of

22 the hippocampus, the dual process view proposes that this structure's impaired efficiency disrupts

23 recollection and other kinds of recall selectively, leaving familiarity intact, and recognition intact to

24 the extent that it is supported by familiarity (see Aggleton & Brown, 1999; Montaldi & Mayes, 2010;

25 Edelstyn, Grange, Ellis & Mayes, 2015). There is also longstanding evidence that amnesic patients'

26 impaired recognition memory improves to the same extent as that of their controls when they are

7

1 given precise elaborative encoding instructions rather than encode in a spontaneous way (e.g.,

2 Mayes, Meudell & Neary, 1980). This is consistent with their usually preserved executive functions

3 and intelligence and strongly suggests that they process and represent informational inputs normally

4 even when their elaborative processing at study and test is self-driven (i.e., spontaneous) rather

5 than guided (see Mayes, 1988 for a review).

6 Unlike the prefrontal cortex / organizational hypothesis, which cannot explain PD familiarity deficits

7 if they are indeed disruptions of a largely automatic kind of memory, the medial temporal

8 lobe/amnesia-like hypothesis can explain a PD familiarity impairment, provided the hypothesis is

9 extended so as to exploit the full neuroanatomical development of the dual process view. The full

10 dual process view also proposes that item familiarity memory is critically mediated by the perirhinal

11 cortex (e.g., Aggleton, et al., 2005; Montaldi, Spencer, Roberts & Mayes 2006; Bowles, Crupi,

12 Mirsattari, Pigott, Parrent & Pruessner, 2007; Montaldi & Mayes, 2010) and its connection to the

13 medial subdivision of the mediodorsal thalamus (Edelstyn, et al., 2015; see also Kaftas & Montaldi,

14 2014 for a review). Therefore, the medial temporal lobe / amnesia-like hypothesis proposes that,

15 although the hippocampus typically becomes dysfunctional first in PD, the perirhinal cortex

16 eventually will do so also typically later in the disease as Braak, Del Tredici, Rüb, de Vos, Jansen Steur

17 & Braak, (2003) have argued. Less commonly, the perirhinal cortex may become dysfunctional at the

18 same time and rate as the hippocampus or, perhaps in rare cases, even earlier so that in unusual

19 cases of PD familiarity may be as impaired as recollection and other kinds of recall or even more

20 impaired (e.g., Davidson, Anaki, Saint-Cyr, Chow & Moscovitch, 2006; Weiermann, Stephan, Kaelin-

21 Lang & Meier, 2010).

22 Recognition and recall memory have been investigated more in PD than have recollection and

23 familiarity, although several more selective studies of these two kinds of memory have been

24 conducted. These studies have usually found that recollection is impaired, but that familiarity is

25 preserved in mild to moderate nondementing PD (Hay, Moscovitch & Levine,2002; Edelstyn *et al.,*

1 2007; Algarabel, *et al.,* 2010; Edelstyn *et al.,* 2010; Shepherd *et al.,* 2013; Rodríguez, Algarabel &

2 Escudero, 2014). However, familiarity has sometimes been found to be impaired as previously

3 indicated (Davidson, et al., 2006; Weiermann, et al., 2010).

4 These findings may differ for two main reasons. First, the differences may have related to how

5 familiarity and recollection were measured. For example, in the Davidson et al. study, it was

6 claimed that familiarity impairments and spared recollection were evident across 3 estimation

7 methods, although in reality, the same data set was used for 2 of these methods (remember-know

8 procedure and word-frequency mirror effect). This is a serious problem given the recognized

9 difficulties with measuring familiarity and recollection accurately and in relatively noise free or

10 unbiased ways, particularly when only small groups are used as they were in Davidson et al.'s study

11 (see Migo *et al.,* 2012). Davidson and his colleagues' third method was the process dissociation

12 procedure. The procedure assumes that the ability to discriminate between intact and recombined

13 word pairs so as to select only intact pairs must depend on recollection alone because it cannot

14 depend on item familiarity. However, it is known that associative familiarity can be found for several

15 kinds of association, including those between words (Bastin, Van der Linden, Schnakers, Montaldi &

16 Mayes, 2010; Harlow, MacKenzie & Donaldson, 2010). If this kind of memory was operating in the

17 Davidson *et al*. study to an appreciable extent, as seems very likely, then the levels of estimated

18 recollection and item familiarity may have been seriously inaccurate given how the process

19 dissociation equations work. This is because the simultaneous equations, related to the inclusion

20 and exclusion conditions of process dissociation procedure, cannot solve three unknowns:

21 recollection, item familiarity and associative familiarity. Indeed, as the conditions were run

22 separately, there may be further unknowns that correspond to the possibly distinct criteria adopted

23 in the two rather different conditions. Second, there may be differences between the patients used

24 in the studies with respect to the stage of their disease, their medication, the pattern of their

25 disease, or specific features such as the severity of PD-related depressive symptoms and executive

26 dysfunction.

1   However, even if all the different findings reflect PD patient-related differences, they are all

2   consistent with the medial temporal lobe/amnesia-like hypothesis provided unknown factors

3   differentially affect the rate at which hippocampal versus perirhinal cortex dysfunction develops.

4   This is not the case for the prefrontal/organizational hypothesis, which cannot explain on its own

5   impaired familiarity and a fortiori familiarity that is more impaired than recollection in PD unless it is

6   argued, contrary to most evidence, that making accurate familiarity judgements is at least as

7   effortful as recollection. It is, of course, possible (indeed likely) that all PD patients show different

8   combinations of prefrontal and medial temporal lobe dysfunction, in which case the possible

9   diversity of familiarity versus recollection deficits indicated above can be explained by the two

10   hypotheses.

11   There is only one PD study, the design of which seems similar to our study reported here, but the

12   findings of which, if reliable, are inconsistent with any combination of the above two hypotheses. In

13   this study, Cohn, Moscovitch & Davidson (2010) argued for a double dissociation between familiarity

14   and recollection as a function of how semantically unrelated word pairs were encoded. When

15   patients encoded the word pairs as they chose, familiarity was impaired and recollection intact,

16   whereas when they generated sentences intended to link the unrelated words together, recollection

17   was impaired and familiarity was intact. However, there are a number of reasons for exercising

18   caution about the reliability of their findings. First, only 9 PD patients and 9 controls were included,

19   when much larger groups are probably needed to provide reliable and replicable results given that

20   the recollection and familiarity deficits are relatively small and PD patients as a group are highly

21   heterogeneous. Second, familiarity and recollection were estimated using a process dissociation

22   procedure. Concerns about the ability of this procedure to provide accurate estimates of familiarity

23   and recollections have already been discussed in relation to earlier work published by the same

24   group (Davidson, et al., 2006). Third, possibly related to the previous problem, control recollection

25   estimates were surprisingly low in the spontaneous encoding condition, and were similar to that of

1   their PD group, as well as a previously studied group of mildly amnesic patients who had undergone

2   unilateral medial temporal lobe resections (Cohn, McAndrews & Moscovitch, 2009).

3   Cohn, *et al.* (2010) argued that the PD recollection deficit suggests hippocampal dysfunction

4   whereas the PD familiarity deficit reflects a strategic (organizational) and related attention deficit

5   caused by striato-frontal dysfunction. However, there is evidence that providing a good semantic

6   elaborative encoding strategy improves PD recall (e.g., Knoke*,* Taylor & Saint-Cyr, 1998; Van

7   Spaendonck*,* Berger, Horstink, Borm & Cools, 1996), and similar effects have been noted in amnesics

8   (e.g., Mayes *et al.,* 1980). Further, even if PD familiarity was increased by giving sentence generation

9   instructions (because it is not completely automatic), contrary to what was reported, it would be

10   expected to increase less than recollection, which is typically more effortful*.* Finally, executive

11   functioning was not assessed. However, it seems likely that, to the extent that they were typical PD

12   patients, it would have been impaired and the patients and their controls would not have performed

13   the unguided sentence generation task in the same way, raising the possibility that the patients may

14   have produced sentences that did not improve recollection.

15   The purpose of the current study was to explore whether memory impairment in PD patients

16   without dementia is better predicted by the prefrontal / organizational hypothesis or by the medial

17   temporal lobe / amnesia-like hypothesis using a design that was not constrained by the limitations

18   identified in the previous Cohn *et al.* (2010) study. To this end, using a large group of patients and

19   controls, we compared the effects of guided elaborative encoding as well as guided elaborative

20   encoding and retrieval versus spontaneous encoding and retrieval on PD and healthy matched

21   control word recognition, familiarity and two kinds of recollection. These kinds of memory were

22   examined using a slightly modified version of the remember/know procedure that fitted well with a

23   source recall task. Executive functions as well as depression were also carefully assessed. The

24   prefrontal / organizational hypothesis predicts that, when guided elaborative encoding (or guided

25   encoding and retrieval) remove the need to use effortful executive processes in order to encode

1 efficiently for subsequent recollection, PD recollection will improve markedly more than that of their

2 controls or possibly even become completely normal. In contrast, the medial temporal lobe /

3 amnesia-like hypothesis predicts that, assuming the piloting was successful, the patients'

4 recollection will improve in the guided conditions relative to the spontaneous encoding condition,

5 but to the same extent as that of the controls. Whether one or both of the hypotheses applies to PD

6 patients, familiarity should not be affected by the two guided conditions. This was because

7 according to the prefrontal / organisational hypothesis, familiarity, depends much less on executive

8 function than intentional recall, and according to the medial temporal lobe / amnesia-like

9 hypothesis, familiarity, is relatively automatic kind of memory. Therefore, this kind of memory

10 should not be affected by elaborative guided encoding designed to enhance semantic links between

11 the paired words and more indirectly the links between the words and thoughts during the more

12 elaborate encoding activity rather than systematically affect the encoding of individual words.

13 By chance, our study's unguided condition (C1) in which encoding was spontaneous and unguided at

14 encoding and retrieval and our partial guidance condition (C2) in which guidance was only provided

15 at encoding were quite similar to the read and sentence generation conditions respectively in the

16 Cohn, *et al.* (2010) study, although the full guidance condition in which guidance was given both at

17 encoding and at retrieval (C3) was unique to our study. However, although both studies required

18 participants to study unrelated word pairs, there was a major difference between C2 and the Cohn

19 *et al.* sentence generation task. Whereas they left participants to generate their own sentences we

20 actually provided sentences known to be beneficial to memory and required participants to judge

21 how well they linked the paired words to ensure that attention was paid to the sentences' meaning.

22 To the extent that executive function was disturbed in the PD patients, one would expect them to be

23 impaired at generating suitable sentences or even generating them at all, and Cohn *et al.* do not

24 report whether this was so. Incidentally, our study allowed us to determine whether anything like

25 Cohn *et al*.'s different encoding-dependent patterns of memory results were produced in a much

26 larger group of PD patients where we could be much more confident that patients really did encode

1   in a similar way to their controls in C2 and C3. We also estimated familiarity and recollection using a

2   modified form of the remember/know (R/K) procedure (Tulving, 1985) rather than the process

3   dissociation procedure (or other procedures for estimating familiarity and recollection) because we

4   believed that the R/K procedure is more reliable if used with care (see Migo *et al.,* 2012 for a

5   discussion of this issue). Our modification also allowed us to measure both objectively scorable

6   source recall of the paired word with which tested words were encoded at study as well as the more

7   standard subjective recollection that scored recall of any other study associates of the tested word.

8   We piloted the three encoding conditions to increase the likelihood that the guided encoding

9   conditions would improve recognition and recollection in healthy older controls.

10   As PD executive function impairments are expected according to the prefrontal / organizational

11   hypothesis we measured some aspects of executive function in our participants and also measured

12   depressive symptoms. As both executive dysfunction and subclinical depressive symptoms were

13   present in the patients, we tested whether they may have contributed to any of the observed

14   memory deficits by correlating them with familiarity and a combined measure of the two kinds of

15   recollection in the three conditions.

16

Materials and method

17

Participants

18

19   Thirty patients were recruited from the Parkinson's disease (PD) outpatient clinic in the Department

20   of Neurology, University Hospital of North Staffordshire. Patients were screened for adverse clinical

21   events or issues (e.g. drastic medication changes, fatigue, distress) that might affect performance in

22   the study. The PD group were in the moderate stages of the condition with a mean illness duration

23   of 6.31 years (SD, 3.34 years), mean medicated modified Hoehn and Yahr disease severity rating of

1   2.53 (SD, 0.9, Hoehn & Yahr, 1967) and mean medicated Unified Parkinson's Disease Rating motor

2   subsection score of 13.38 (SD, 5.08, Fahn & Elton, 1987).

3   A group of twenty-two healthy controls provided control data for the recognition memory tasks.

4   They were matched to the PD patients for age and current levels of functioning (*Mini-mental state*

5   *examination,* Folstein*,* Folstein & McHugh, 1975; *Wechsler Abbreviated Scale of Intelligence*,

6   Wechsler, 1999).

7   The demographic, neuropsychological and clinical (patients only) characteristics of the healthy

8   control and PD groups are provided in Table 1.

9

10

11   Table 1 around here

12

13

14   Patients were tested in a medicated state (within 90 minutes of taking their morning medication),

15   and at the time of testing, were on a mixture of medication regimens that included either l-dopa, a

16   second generation dopamine agonist (pramipexole, ropinirole or rotigotine) or a monoamine B

17   enzyme inhibitor as either monotherapy or in various combinations. The mean l-dopa equivalent

18   dose was 635.71 mg (SD, 463.86).

19   Exclusion criteria for all participants included a *Minimental* score of 25 or less, presence or a history

20   of a psychiatric or neurological illness including diagnosable dementias such as Alzheimer's Disease

21   (apart from the PD patients in the index group), history of substance abuse (such as alcoholism),

22   currently taking antidepressants, learning difficulty (including dyslexia), or English as a second

23   language. Additional exclusion criteria for the patients included a history of visual hallucinations

1 and/or delusions, dyskinesias, impulse control disorders, and commencement of dopaminergic

2 medication within the two months prior to entering the study.

3 The study was approved by the Keele University Faculty of Humanities and Social Sciences Research

4 Committee and South Staffordshire NHS Research Ethics Committee, and conducted in accordance

5 with Good Clinical Practice.

6 ***Experimental Recognition Memory Test***

7 ***Stimuli***   Three Yes-No recognition memory tests (RMT) were constructed from a pool of 150

8 verbal paired associates (VPAs). The VPAs were created by asking fifteen undergraduates to

9 generate an associate for each of one hundred and fifty words (mean imagery 6.05, SD 0.49; mean

10 concreteness 6.27, SD 0.65; mean frequency 3.39, SD 0.90). The VPAs that were least frequently

11 associated to the first word formed the second word in each word pair. This process created further

12 one hundred and fifty weakly associated VPA. The VPAs were randomly assigned to the three

13 recognition memory tests, with each version containing fifty word pairs. In the spontaneous

14 encoding and retrieval condition, the target stimuli comprised word pairs (e.g. **IRON - CREASE**).

15 A further one hundred and fifty nouns (mean imagery 5.52, SD 0.68; mean concreteness 5.33, SD,

16 1.13, mean frequency 3.28, SD, 1.34) formed the lures (i.e. new words) for the test sessions. None

17 of the lures appeared in the target VPAs used during the study phase. The distracters were

18 randomly assigned to three packs, each containing fifty words. At test, first or second words from

19 the VPAs and never both words from any one pair were randomly selected and presented in a

20 random mixed order, intermixed with fifty lures. Assignment of packs of lures and target VPAs, and

21 whether the first or second word from each word pair was presented at test, were counterbalanced

22 across participants.

23 ***Procedure***   All participants completed the same three RMT conditions, administered in the same

24 sequence (C1, followed by C2, in turn, followed by C3). The sequence in which the conditions were

1 completed was not counterbalanced to prevent carry-over of organisational strategies between C3

2 and C2, and between each guidance condition and the baseline / no guidance condition (C1). Each

3 RMT was completed at the same time of day on three non-consecutive days. Each condition was

4 presented and delivered in exactly the same manner in all respects apart from the form in which the

5 target VPAs were presented at study and the accompanying instructions provided at study and test.

6 In the first RMT condition, C1, participants viewed the VPA and were given standard encoding and

7 retrieval instructions (see the following section for details)*. In the second and third RMT conditions,

8 C2 and C3, the VPA were embedded in sentences that provided a connection between the two

9 associates. For example, for the VPA '**TELEPHONE** – **PLUG**', the target sentence read "The

10 **TELEPHONE** was located close to the wall **PLUG**". Directed encoding guidance was combined with

11 standard retrieval instructions in C2, and with directed retrieval guidance in C3.

12 The target VPAs in C1, and sentences in C2 and C3, were each displayed for 9000 msec during the

13 *study* phase, and participants also had a 9000 msec window in which to either endorse or reject the

14 probe during the test phase. The second test stage for endorsed items which required participants

15 to make a *remember, know* and source recall response was not time constrained.

16 A 25 minute interval that separated the *study* and *test* phases was filled with non-verbal

17 neuropsychological testing (see below). The session occurred at the same time in the morning for all

18 of the participants. Prior to administering the first RMT, participants were familiarized with the

19 experimental set-up, stimuli and task requirements, including types of responses they were required

20 to make using tailored practice tests. None of the word pairs or new words used in the practice test

21 appeared as new words or word pairs in the main RMT.

22 ***Instructions***   The following "standard" instructions were given to participants prior to

23 commencing the study phase in C1: *'I am going to show you fifty pairs of words written in bold*

24 *letters that are slightly related to each other. Read each word pair aloud. Do your best to commit*

16

1   *them to memory because I will be testing your memory for the items later'*. The "standard" retrieval

2   instructions provided immediately prior to test in C1 and C2 were: "*Now, I am going to test your*

3   *memory for the words I have just shown you."*

4   The guided encoding instructions in C2 and C3 were: '*I will show you fifty pairs of words in bold*

5   *letters. Each will appear in a sentence that helps connect the words. I want you to read the sentence*

6   *aloud and say whether you think that it relates the words well or not. This will help you remember*

7   *the words when I test your memory shortly after the study session has finished. We have tried to*

8   *make the sentences so that they help relate the words so do not worry if you think that most or, even*

9   *all, the sentences do this.'*

10   The additional guidance provided at test in C3 was '*It will help you to do the task if you try to*

11   *remember the sentence that you were shown during the study session.'*

12   ***Performance Measures***   Correct identification of a target item was defined as a *hit*, whilst

13   false recognition of a lure was termed a *false alarm*. Following each endorsement, irrespective of

14   whether it was a hit or false alarm, participants were **first** asked to make a judgement about

15   whether their recognition was accompanied by either recollection of specific details, in addition to

16   the word paired with the tested word at study associated with studying the tested word earlier

17   ('*remember'* response) or by feelings of familiarity without any recollection ('*know'* response). Recall

18   of the source word was prompted by the experimenter if it had not already been spontaneously

19   produced.

20   A correction has been made to the data to eliminate extreme scores in accordance with Snodgrass

21   and Corwin's (1988) recommendation. Familiarity and recollection memory discrimination scores

22   were made by first computing the familiarity and recollection hit rates and false alarm rates. In order

23   to calculate the hit and false alarm rates for familiarity, it was assumed that recollection and

24   familiarity are stochastically independent at retrieval, and therefore, Yonelinas and Jacoby's (1995)

1   independence formula has been applied to the corrected *know* scores (Familiarity = *know*/ [1-

2   *remember*]). Estimates of recognition and familiarity were calculated using signal detection theory

3   (*d'*), and a threshold measure (*pr*) is reported for subjective recollection and source recall (i.e. hit

4   rate minus false alarm rate). Based on fMRI data, introspective experience and general plausibility

5   arguments, there are strong reasons to suppose that this assumption is much nearer to reality than

6   that familiarity and recollection have an exclusivity relationship or that, every recollection response

7   is always accompanied by a familiarity response (redundancy) (see Migo, Mayes & Montaldi, 2012

8   for a full development of these arguments).

9   ***Remember-know instructions***   Participants were instructed that a *remember* response

10   could be given only if, when presented with a probe item, they recollected at least one of the

11   following*:* (i) the item that appeared just before or after the currently being tested "probe" item

12   during the study session*;* (ii) personal memories, mental images, or words that came to mind when

13   the probe was presented during the study session; (iii) an emotional reaction that the currently

14   being tested probe triggered during the study session. These instructions aim to ensure that

15   recollection is only reported when a participant has recalled one or more things that they were

16   thinking of when they were processing the probe during study. It is important to stress that

17   remember judgements specifically *did not* include source recall of the word paired at study with the

18   tested word.

19   *Know* responses were only recorded if participants recognised the probe as having been presented

20   in the study session, but did not recall any specific details, including the paired word about it from

21   the study session. Although guess responses are sometimes also included in the *remember/know*

22   procedure, they were not used for two reasons. Their use is unwarranted because familiarity

23   memory discrimination scores were corrected using the familiarity false alarm rate so if any

24   familiarity response was really a guess rather than a weak familiarity response, this was fully

25   corrected. The second reason that we did not use guess responses was that it is likely that the extra

1 complexity will confuse participants. Such confusion is highly undesirable because there is evidence

2 that unless instructions are kept simple and are fully understood, participants can too frequently fail

3 to follow them properly (see Migo *et al.,* 2012 for a full discussion of these points).

4 Participants were asked to justify each *remember* and *know* judgement throughout the test phase to

5 ensure that they maintained a full understanding of the criteria for making these types of decisions

6 in line with published recommendations on measuring recollection and familiarity using the

7 *remember/know* procedure recommended by Migo *et al.* (2012).

Neuropsychological Tests

9 Controls and patients completed a depression questionnaire (*Hamilton Depression Inventory,*

10 Reynolds & Kobak, 1995), a test of executive function (*The Brixton Test of Spatial Anticipation,*

11 Burgess & Shallice, 1997) and an independent measure of delayed verbal recall (*Logical Memory*

12 delayed verbal recall subtest from *The Wechsler Memory Scales,* Wechsler, 1997), which previous

13 studies indicate are impaired in PD (e.g., Cooper, Sagar, Jordan, Harvey & Sullivan, 1990; Crescentini,

14 Mondolo, Biasutti & Shallice, 2008; Edelstyn *et al.,* 2007; Shepherd *et al.,* 2013). The questionnaire

15 and both tests were administered according to their respective manuals.

Data Analysis

17 To examine the effects of guidance instructions on the performance measures, a series of 2 by 3

18 mixed analyses of variance (ANOVA) were conducted with Group (PD group vs healthy controls) as

19 the between subjects factor; and memory condition (C1 vs C2 vs C3) as the within subjects factor.

20 Significant main effects and interactions were investigated further using planned pair-wise

21 comparisons.

22 The effect of increasing strategic guidance on memory performance within each group was also

23 examined using paired samples t-tests.

1   A series of bivariate correlational analyses, using Pearson's product moment coefficients, explored

2   the relationship between the measures of subclinical depressive symptoms, executive function,

3   delayed verbal recall, RMT performance measures (familiarity [d'] and a composite measure of

4   recollection [pr] in C1, C2 and C3) in a subset of participants (17 PD patients and 10 healthy

5   controls). The reported correlations have not been corrected for multiple comparisons as their

6   purpose was to examine whether there was any evidence, however, weak, that impaired

7   dysexecutive processing driven by prefrontal dysfunction and/or subclinical depression might also be

8   contributing, at least to a small extent to the PD memory deficits.

9

10 **Results**

11 The raw hit and false alarm rate (HR, FAR, respectively) means and standard deviations for item

12 recognition memory (RM), *know, remember* and source recall by group are shown in Table 2. The

13 estimates of item RM (*d'*), familiarity (*d'*), subjective recollection (*pr*) and source recall (*pr*) are also

14 shown in Table 2 and in Figure 1.

15

16

17   Table 2 and Figure 1 around here please

18

19

20 ***Comparisons between PD patients and Controls***   Analysis of item recognition *(d')* revealed

21 main effects of *Group* ($F(1,50) = 12.81$, $\eta^2 = .23$, $p = .001$) and *Condition* ($F(2,50) = 3.83$, $\eta^2 = .08$, $p = $

22 .026), but the *Interaction* between *Group* and *Condition* was not significant ($F(2,50) = 0.78$, $\eta^2 = .02$,

1   *p* = .46).   Item recognition in the PD patients was significantly lower than in the healthy controls in

2   C1 (*t*(50) = -3.53, *p* = .001) C2 (*t*(50) = -2.34, *p* = .02) and C3 (*t*(44) = -2.85, *p* = .01).

3   The second ANOVA of familiarity *(d')* showed a main effect of *Group* (F(1,50) = 4.01, η^2 = .0.84, *p* =

4   .051) but not of *Condition* (F(2,50) = 2.38, η^2 = .05, p =. 9), and the interaction was also not significant

5   (F(2,50) = 2.00, η^2 = .04, *p* =. 14). Familiarity in the PD patients was significantly lower than in the

6   healthy controls in C1 (*t*(50) = -2.18, *p* = .034) but not in either C2 (*t*(50) = -1.39, *p* = .17), or C3 (*t*(44)

7   = -0.83, *p* = .39) where partial and full guidance were provided respectively.

8   Analysis of subjective recollection *(pr)* was marked by main effects of *Group* (F(1,50) = 27.46, η^2 =

9   .38, *p* <. 001) and *Condition* (F(2,50) = 17.50, η^2 = .29, *p* < .001), but the interaction was not

10   significant (F(2,50) = 1.69, η^2 = .04, *p* =. 19). Subjective recollection in the PD patients was

11   significantly lower than in the healthy controls in each of the three conditions (C1: *t*(50) = -6.21, *p*

12   <.001; C2: *t*(50) = -2.99, *p* = .004 and C3: *t*(44) = -4.62, *p* < .001).

13   The final ANOVA of source recall *(pr)* showed main effects of *Group* (F(1,50) = 7.58, η^2 = .16, *p* =

14   .009) and *Condition* (F(2,50) = 6.14, η^2 = .14, *p* = .003), but the interaction was not significant

15   (F(2,50) = 7.03, η^2= .03, *p* = .30). Source recall in the PD patients was significantly lower than in the

16   healthy controls in C1 and C3 (C1: *t*(50) = -2.44, *p* = .02; C3: *t*(39) = -2.35, *p* = .02) but not in C2 (*t*(50)

17   = -0.41, *p* = .69).

18   To compare the relative severity of impairment, estimates of familiarity, subjective recollection and

19   source recall in the unguided/spontaneous encoding and retrieval condition (C1) were converted to

20   standard scores. A paired-samples t-test showed familiarity (z = -0.32) to be significantly less

21   impaired than both subjective recollection (z= -1.45,   *t* (29) =10.0, p <.001) and source recall (z = -

22   1.65,   *t* (29) = 5.83, p <.001), whereas subjective recollection and source recall showed comparable

23   levels of decline (*t* (29) = 0.96, p = .34).

1 ***Effects of Guidance Within Each Group***   Paired sample t-tests showed PD patients' RM,

2 subjective recollection and source recall each improved with guidance (C2, C3) compared to baseline

3 (C1). But the amount of memory improvement did not increase when guidance was given both at

4 retrieval and encoding (C3) compared to encoding alone (C2) (RM, C1-C2: $t(29)$ = -2.8, p = .009; C1-

5 C3: $t(29)$ = -2.92, p = .007; C2-C3: $t(29)$ = -0.28, p = .78; subjective recollection, C1-C2: $t(29)$ = -5.12, p

6 < .001; C1-C3: $t(29)$ = -5.0, p < .001; C2-C3: t(29)=-1.14, p = .26; source recall, C1-C2: $t(29)$ = -2.7, p =

7 .011; C1-C3: $t(29)$ = -2.69, p = .01; C2-C3: $t(29)$ = -0.37, p = .71).

8 In contrast, there was no effect of guidance on PD familiarity between any of the contrasted

9 guidance conditions (C1-C2: $t(29)$ = .16, p = .88; C1-C3: $t(29)$ = .3, p = .77; C2-C3: $t(29)$ = .08, p = .93).

10 The healthy controls showed improvements in both subjective recollection and source recall

11 between the unguided (C1) and full guidance (C3) conditions (subjective recollection, $t(15)$ = -3.1, p <

12 .01 and source recall: $t(10)$ = -2.51, p = .03, respectively); partial and full guidance conditions C2-C3

13 (subjective recollection: $t(15)$ = -2.55, p = .02; source recall: $t(10)$ = -2.23, p = .05) but not between

14 unguided (C1) and partial guidance (C2) (subjective recollection: $t(21)$ = .78, p = .44; source recall:

15 $t(22)$ = -.36, p = .72)

16 A borderline decrease in familiarity was present between unguided and partial guidance conditions

17 ($t(21)$ = 2.04, p = .06) but not between unguided and full guidance or between partial and full

18 guidance (C1-C3: $t(15)$ = 1.94, p = .072; C2-C3: $t(15)$ = 1.24, p =. 23).

19 There were no changes in RM between any of the guidance conditions (RM, C1-C2: $t(21)$ = -0.49, p =

20 .63; C1-C3: $t(15)$ = -.85, p = .41; C2-C3: $t(15)$ = -.38, p = .64.

21 In summary, patients with mild to moderate PD (mean HY 2.53) displayed deficits in source recall

22 and subjective recollection whereas item familiarity was markedly less impaired. There was no

23 evidence that the PD group gained more advantage from partial guidance (at study only) than the

24 controls (compare C1 with C2). However, patient subjective recollection and source recall remained

22

1 impaired even with full guidance at test as well as at study, and, critically, the two groups improved

2 to the same extent when C3 was compared to C1. These findings are illustrated in Figure 2.

3

4

5 Figure 2 around here please

6

7

8 ***Neuropsychological and Depression Scores***

9 Scores on the Brixton Test, Logical Memory and Hamilton Depression Inventory are shown in Table

10 3.

11

12 Table 3 around here, please

13

14 Compared to the healthy controls, PD patients showed evidence of executive dysfunction (*Brixton*

15 *Test*, *t* (25) = -2.6, p = .02), impairment in delayed verbal recall (*Logical Memory* delayed verbal

16 recall, *t* (25) = -5.07 p <.001) and significantly elevated rates of subclinical depressive symptoms

17 (*Hamilton Depression Inventory [HDI], t* (25) = -2.45 p =.021).

18 **The PD Group**   HDI scores weakly correlated with overall recollection in the unguided,

19 partial and full guidance conditions, respectively (C1: *r*(17) = -0.36, *p* = .077; C2: *r*(17) = - 0.46, *p* =

20 .03; C3: *r* (17) = -.36, *p* = .078, respectively). HDI scores also failed to correlate with the C3-C1

21 difference score for overall recollection (r=-0.04, p=0.83).

1   Delayed story recall correlated with overall recollection in each of the three conditions (C1: r (17) =

2   0.62, p = .004; C2: r (17) = 0.59, p = .007 and C3: r(17) = -0.48, p = .024).

3   Executive function was correlated with unguided overall recollection rates in C1 only (r (17) = 0.46, p

4   = .03), and failed to correlate with overall recollection in the presence of partial guidance at

5   encoding (C2: r (17) = 0.31, p = .11) and full guidance at encoding and retrieval (C3: r (17) = 0.13, p =

6   .31). Executive function also failed to correlate with the C3-C1 difference score for overall

7   recollection (r=-0.11, p=0.64).

8   Familiarity rates in the unguided and guided conditions failed to correlate with HDI scores (C1: r (17)

9   = -0.07, p = .39; C2: r (17) = 0.14, p = .29; C3: r (7) = 0.09, p = .37), executive function (C1: r (17) =

10   0.27, p = .14; C2: r (17) = -0.34, p = .09; C3: r (17) = -0.08, p = .38), delayed recall (C1: r (17) = 0.27, p

11   = .15; C2: r (17) = -1.02, p = .35; C3: r (17) = -0.04, p = .44) and overall recollection (C1: r (17) = 0.26,

12   p = .15; C2: r (17) = -0.20, p = .23; C3: r (17) = -0.22, p = .20).

13   Finally, HDI scores showed border-line correlations with delayed recall and executive function (r (21)

14   = -0.39, p = .06; r (21) = -0.41, p = .05, respectively).

15   In summary, PD executive function correlated with unguided overall recollection only, and failed to

16   correlate with the C3-C1 difference score for overall recollection. HDI scores weakly correlated with

17   overall recollection in the unguided, partial guided and fully guided conditions, but failed to

18   correlate with the C3-C1 difference score for overall recollection..

19   **The Healthy Controls**   Overall recollection failed to correlate with HDI scores and

20   familiarity estimates and in the unguided and guided conditions (depression, C1: r (10) = 0.28, p =

21   .22; C2: r (10) = 0.12, p = .37; C3: r (10) = -0.38, p = .14; familiarity,C1: r (10) = 0.03, p = .46; C2: r (10)

22   = 0.47, p = .09; C3: r (10) = 0.42, p = .12). Executive function was not correlated to overall

23   recollection in any of the conditions (C1: r (10) = -0.23, p = .26; C2: r (10) = 0.19, p = .30; C3: r (10) = -

1   0.20, p = .29). Delayed recall weakly correlated to overall recollection in C2 but not in either C1 or C3

2   (C1: r (10) = 0.14, p = .35; C2: r (10) = -0.52, p = .06; C3: r (10) = -0.36, p = .16).

3   Estimates of familiarity correlated to executive function and delayed recall in the full guidance

4   condition only (executive function, C1: r (10) = 0.34, p = .17; C2: r (10) = -0.21 , p = .28; C3: r (10) =

5   -0.6, p = .034; delayed recall, C1: r (10) = 0.23, p = .26; C2: r (10) = 0.07, p = .4; C3: r (10) = 0.05, p =

6   .05). There were no correlations between familiarity and depression rates (C1: r (10) = -0.33, p = .18;

7   C2: r (10) = 0.04, p = .46; C3: r (10) = -0.06, p = .43). There were also no correlations between

8   depression levels and executive function (r (10) = -0.33, p = .18), depression levels and delayed recall

9   (r (10) = 0.02, p = .48), or executive function and delayed recall (r (10) = -0.36, p = .16).

10

11   **General discussion**

12   In all three of the learning and test conditions used in our study, a large group of non-dementing

13   mild to moderate PD patients were impaired at the two kinds of cued recall examined: source recall

14   of words paired at study with the tested words and subjective recollection of other associations with

15   the tested words from the study episode. Contrary to the findings in our previous work, the patients

16   were also impaired at word familiarity, although their familiarity deficit was very modest and

17   significantly smaller than their impairments in both kinds of cued recall. Given the dependence of

18   recognition test performance on both familiarity and recollection, it was unsurprising that the

19   patients showed an overall deficit in word recognition in each of the 3 conditions. In separate tests,

20   the patients also were clearly impaired at delayed free recall of short stories (i.e., delayed logical

21   memory) and, as is often found with mild to moderate PD patients, they showed impaired executive

22   functions as well as significantly elevated levels of subclinical depressive symptoms relative to their

23   controls.

1   Most importantly, in the PD patients as well as their controls, both kinds of cued recall were

2   significantly increased with full guidance at encoding and retrieval compared to the unguided

3   condition(C3 versus C1). As the amount of cued recall improvement shown by patients and controls

4   was similar, this indicates that full guidance relative to no guidance for both groups was equally

5   beneficial for their cued recall. This finding fits broadly with what should be expected if the PD

6   patients' cued-recall impairment is primarily driven by a medial temporal lobe / amnesia-like deficit

7   rather than a breakdown in prefrontal cortex-dependent organisational processes.

8   However, although there was no cued recall interaction between group and degree of guidance,

9   individual t-tests suggested that PD patients' main cued recall benefit was provided by encoding

10   guidance and that they gained little extra from additional retrieval guidance. In contrast, with

11   controls, these effects were the other way round with them gaining little from encoding guidance,

12   but significantly from retrieval guidance. There is a danger of overinterpreting this effect, which

13   may be a statistical anomaly and can at most only indicate a weak effect. Such a weak effect may or

14   may not turn out to be genuine when examined by a future study that would have to include many

15   more patients than did the current study (probably around 100) to have sufficient power for a

16   convincing examination. If it did prove to be a real but weak effect, it would most likely indicate that

17   patients and controls used a slightly different strategy with the guidance offered. For example, the

18   less confident patients might have tried to think of what they had encoded when tested even when

19   not given explicit retrieval instructions to do this whereas the more confident controls did this much

20   less unless explicitly asked to do so in the full guidance condition. Whether or not this happened,

21   however, the fact that full guidance improved patient and control cued recall to the same significant

22   degree, is incompatible with the PD recollection deficit being appreciably affected by a prefrontal

23   /executive impairment. PD word familiarity, on the other hand, did not differ across the 3

24   conditions, being unaffected by the kinds of guidance which we provided. Although the group by

25   guidance condition was not significant, the controls showed an insignificant tendency to perform

26   more poorly on this measure in the guided conditions (C2 and C3) than in the spontaneous condition

26

1   (C1), which accounted for the hint of a PD familiarity deficit in the spontaneous versus guided

2   conditions. However, the lack of a guidance condition interaction with group for familiarity together

3   with the lack of effect of condition indicates that the weakly significant PD familiarity deficit noted in

4   C1, if reliable, is more likely to reflect perirhinal cortex/medial temporal cortex dysfunction rather

5   than frontal cortex dysfunction, contrary to the proposal of Cohn et al. (2010). This point has been

6   discussed previously in the Introduction in relation to functional imaging work (see Henson et al.,

7   1999), prefrontal lesion studies (Duarte et al., 2005; MacPherson, et al., 2008) and behavioural

8   evidence from older adults (Anderson, et al., 2008), and is explored further in later sections of the

9   Discussion with reference to modulation of hippocampal function by the dopaminergic mid brain,

10   evidence of hippocampal atrophy in nondementing PD, and Braak *et al.*'s (2003) staging model of

11   PD.

12   The significant correlation of delayed free recall of short stories with the measure of overall

13   recollection, a composite measure of both kinds of cued-recall, but not word familiarity, in all three

14   conditions was consistent with the view that these kinds of cued-recall memory have overlapping

15   functional and neural mechanisms whereas they have relatively distinct underlying processes and

16   neural bases from item familiarity. This view is further reinforced by the failure of familiarity to

17   correlate with overall recollection in guided as well as unguided conditions.

18   The significant correlation of executive functions in the PD patients only with unguided overall

19   recollection in C1 suggests that the patients' impaired executive functions may have slightly

20   worsened their spontaneous processing, but that the guided conditions markedly reduced the need

21   to rely on these dysexecutive functions. The patients' executive function failed to correlate with the

22   C3-C1 difference score for overall recollection which is again consistent with the view that a frontal

23   dysexecutive impairment is not a major cause of the PD memory disorder. Finally, the absence of a

24   correlation between the patients' executive functions and familiarity in any of the conditions is

25   compatible with familiarity being unhelped by guidance. This finding is expected because familiarity

1   seems to be a relatively automatic kind of memory, which is dependent on the perirhinal cortex and

2   its connections. However, weak and, as yet unspecified, prefrontal contributions may be present

3   that need not involve the frontal executive processes that probably support intentional cued-

4   recall. Even if such executive processes are involved, our results suggest that familiarity depends on

5   them much less than the typically more demanding and effortful cued recall memory.

6   The tendency for subclinical depressive symptoms in the PD patients to correlate with overall

7   recollection, but not word familiarity, in all three conditions, is consistent with subclinical depressive

8   symptoms contributing to their recollection deficit but not their poorer familiarity memory.

9   This pattern of results is broadly what would be expected if hippocampal dysfunction/degeneration

10   was a major contributor to our patients' impaired recollection and a smaller degree of

11   dysfunction/degeneration in the perirhinal cortex was a major contributor to their slight impairment

12   in word familiarity. In other words, our findings are broadly consistent with the medial temporal

13   lobe/amnesia-like hypothesis of PD memory. Hippocampal dysfunction would, of course, give rise to

14   inefficient working of a more extended neural system that mediates the kind(s) of memory that

15   underlie recall, whereas perirhinal cortex dysfunction would give rise to inefficient working of a

16   distinct, more extended system that mediates the kind(s) of memory that underlie familiarity.

17   This interpretation of our findings is consistent with evidence that dopaminergic inputs from the

18   dopaminergic midbrain ventral tegmental area modulate activity in the hippocampus and perirhinal

19   cortex as well as the parahippocampal and entorhinal cortices within the medial temporal lobes

20   through feedback circuits involving projections from the hippocampus via the nucleus accumbens

21   and perirhinal cortex respectively (see Lisman, Grace & Duzel, 2011 for a review). The role of

22   hippocampal dysfunction in the PD recollection and recall impairment is specifically supported by

23   rodent and imaging studies of healthy human adults, which show that the hippocampus and the

24   ventral tegmental area form a functional loop controlling the entry of novel and salient/goal-

25   directed information into long-term memory (e.g., Bunzeck, *et al.,* 2007; Gasbarri, Sulli & Packard,

1   1997; Bethus, Tse & Morris, 2010; Chowdhury, Guitart-Masip, Bunzeck, Dolan & Duzel, 2012; for a

2   review see Lisman & Grace, 2005). Dopamine D1 and D2 receptors are also implicated in the

3   persistence / slow consolidation of hippocampal-dependent memories (Laszy, Laszlovsky &

4   Gyertyan, 2005; Hammad & Wagner, 2006; O'Carroll, Martin, Sandin, Frenguelli & Morris, 2006;

5   Takahashi, *et al.,* 2008; Lisman, *et al.,* 2011). However, although functional disruption of the

6   hippocampal recall-related circuits will result from ventral tegmental area degeneration even if the

7   hippocampus remains largely structurally intact, there is also evidence that indicates there is

8   hippocampal neuropathology in nondementing PD with volume loss particularly associated with the

9   CA2-4 subfields/dentate gyrus (Pereria *et al.,* 2013). Volumetric imaging studies in nondementing PD

10   also report an association between recall but not recognition and hippocampal neuropathology (e.g.,

11   Laakso, *et al.,* 1996; Riekkinen, Kejonen, Laakso, Soininen, Partanen & Riekkinen, 1998; Junqué, *et*

12   *al.,* 2005; Bouchard, *et al.,* 2008; Ibbarretxe-Bilbao, *et al.,* 2008; Carlesimo, *P*iras, Assogna, Pontieri,

13   Caltagirone & Spalletta, 2012; Filoteo, Reed, Litvan & Harrington, 2014; however, for a counter

14   argument see Camicoli, Moore, Kinney, Corbridge, Glassberg & Kaye, 2003; Nagano-Saito, et al.,

15   2005; Tam, Burton, McKeith, Burn & O'Brien, 2005; Apostolova, *et al.,* 2010).

16   However, the overall pattern of our data (for example, the statistically weaker correlational

17   evidence, which should be taken as suggestive rather than conclusive) indicates that our main

18   findings do not eliminate the possibility that there is a weak contribution from impaired

19   dysexecutive processing driven by prefrontal dysfunction. Furthermore, if it occurs, the extent of

20   prefrontal contribution is likely to vary across different PD patients, although the factors underlying

21   this remain to be fully clarified so a significant effect may not always be found except within very

22   large groups of patients.

23   Our finding that subclinical depressive symptoms weakly correlated with overall recollection, but not

24   familiarity, in all three conditions suggests that depression in PD may impair recollection selectively

25   in a way that does not need to depend strongly on its effect on executive functions. This is what

1 would be expected if depression causes an amnesia-like memory problem by disrupting the

2 hippocampus because this should impair recollection/source recall equally regardless of whether

3 processing is spontaneous or guided so as to depend less on frontal executive functions, i.e., in all

4 three conditions. In contrast, if depression also acts on the frontal cortex to disrupt executive

5 functions, then the strongest effect should be on the spontaneous condition because this depends

6 more on these functions. This suggests that, if depression also disrupts overall recollection via its

7 disruptive effect on executive functions, then it should correlate with the difference in overall

8 recollection between the spontaneous and most guided condition, i.e., C3 and C31. However, it did

9 not, which also suggests that depression primarily disrupts overall recollection via its effect on the

10 hippocampus to produce an amnesia-like condition, although all these correlations were weak so

11 future work should use much larger participant numbers.

12 It is well-established that prolonged depression is particularly associated with hippocampal atrophy,

13 which is believed to play a key role in the memory disorder found in depression (e.g., Gradin & Pom,

14 2008). An interesting, if speculative, possibility is that a component of this atrophy is decreased adult

15 neurogenesis in the dentate gyrus of the hippocampal (for reviews, see Kempermann & Kronenberg,

16 2003; Sahay & Hen, 2007). Such decreased neurogenesis in the anterior dentate gyrus would

17 negatively disrupt cellular processes underlying pattern separation in the CA1-4/dentate gyrus

18 subfields of the hippocampus (e.g., Sahay, *et al.,* 2011). . This kind of processing is believed to

19 underlie the kind of associative memory that supports cued recall (e.g., Clelland, *et al.,* 2009).

20 There is also evidence for less marked and more delayed degeneration in the perirhinal cortex in PD,

21 which is predicted by the staging model according to which neuropathology emerges in the CA2

22 fields of the hippocampus before it extends to the medial portion of perirhinal cortex (Braak *et al.,*

23 2003, see also Braak *et al.,* 2006; Braak & Del Tredici, 2008; see also Pereira, *et al.,* 2013). However,

24 the relationship of the staging of these degenerative changes and their associated memory changes

25 to clinical severity needs further exploration. In particular, the factors that underlie the relative

1   rates at which hippocampal and perirhinal cortex degeneration occur and the precise mechanisms

2   underlying the degeneration are currently unknown.

3   Interestingly, if perirhinal cortex atrophy is mainly responsible for our patients' marginal decline in

4   familiarity this may help explain why we have not previously found this kind of memory to be

5   impaired in mild to moderate PD. Unlike our other studies (Edelstyn *et al.,* 2007; Edelstyn *et al.,*

6   2010; Shepherd *et al.,* 2013), in this one, before testing memory, we used a 25 minute delay during

7   which participants were occupied with other tasks. This would have produced interference, which,

8   according to Sadeh, Ozubko, Winocur & Moscovitch (2014), is the main mechanism responsible for

9   forgetting of perirhinally-supported familiarity memory. This suggestion is consistent with the two

10   previous PD studies reporting a selective familiarity deficit, where a filled delay of 10 minutes

11   (Davidson, *et al.,* 2006) and 30 minutes (Weiermann, *et al.,* 2010) was introduced between study

12   and test. In addition, it has been argued that, following perirhinal cortex damage, object/item inputs

13   can no longer be integrated at the highest level in the ventral stream so that item representations

14   are likely to become more similar to each other. As interference is a similarity-based process, it will

15   increase following perirhinal cortex damage so that familiarity impairments will be greater after a

16   delay (Saksida & Bussey, 2010).

17   Our findings do not preclude the possibility of a PD-related impairment in executive functions

18   contributing to our patients' verbal memory disorder, but they were only very weakly supportive of

19   the possibility. The patients' executive function scores did show a tendency to correlate with overall

20   recollection only in the unguided, spontaneous condition (C1), but if this correlation reflected a

21   causal influence on recollection of the patients' executive impairment, they should have gained

22   more from guidance, particularly full guidance (C3) than their controls. But the interactions between

23   both kinds of recollection and condition were not significant. Recollection/recall did improve more in

24   the patients between C1 and C2, but, this effect was reversed between C2 and C3 with control

25   recollection tending to improve more than patient recollection, albeit not significantly so. This

31

1   finding is consistent with a recent PD study reporting executive function to be weakly related to

2   verbal episodic recall using factor analysis, canonical regression and structural equation modelling

3   (Alonso Recio, Martin, Carvajal, Ruiz & Serrano, 2013)

4   Future research on different kinds of guidance may clarify whether PD recall memory is

5   disproportionately improved by any kinds of guidance, but our evidence is not suggestive that it

6   does. This would be expected in so far as PD disrupts the functioning of frontal regions that mediate

7   executive functions that facilitate processing at encoding and retrieval. PD has long been associated

8   with dysfunction in some frontal regions and there is evidence that early non-demented PD patients

9   show prefrontal cortex atrophy (e.g., Bruck, Kurki, Kaasinen, Vahlberg & Rinne, 2004). Although

10   whether the frontal impairments found in PD impair executive functions that disrupt recall remains

11   to be convincingly shown, depression is known to disrupt executive functions and to impair recall

12   (e.g., see Channon & Green, 1999) as well as to disrupt frontal functioning (e.g., Baxter, *et al.,* 1989)

13   so PD patients with subclinical depressive symptoms may well suffer from executive function deficits

14   that exacerbate their amnesia-like recall deficits that are caused by hippocampal dysfunction. In so

15   far as subclinical depressive symptoms contributes to both prefrontal and hippocampal dysfunction,

16   resolution would require a very large study that uses a regression analysis or the use of structural

17   and possibly functional MRI in quite a large study to discover how strongly each structural region

18   relates to the recollection deficit.

19   Even though our study may seem, on the surface, similar to that of Cohn *et al.* (2010), particularly

20   with respect to our conditions C1 and C2 and their spontaneous ("Read-only") and guided

21   ("Sentence generation") conditions, the fact that they used process dissociation procedure meant

22   that they tested half the word pairs their participants encoded with a word recognition test and half

23   with an associative word recognition test. Our participants, in contrast, were only tested on word

24   recognition and had to recall the paired words rather than recognize them. Also, Cohn and her

25   colleagues did not set out to deliberately test the hippocampal/amnesia-like hypothesis against the

1 prefrontal/organizational hypothesis. The failure of the sentence generation task to improve

2 recollection in Cohn *et al.'s* patients unlike in their controls may have arisen because their patients'

3 presumed executive deficits led to their not generating suitable sentences. The need to do this was

4 obviated in our study by the provision of appropriate sentences. However, this difference does not

5 explain the very poor estimated recollection scores of Cohn *et al.'s* controls in their Read condition,

6 which led to the apparently normal recollection scores of their patients in this condition. We believe

7 it to be more likely that this strange finding as well as Cohn *et al.'s* findings with word familiarity

8 arose because of distortions resulting from their use of the process dissociation procedure. In

9 particular, associative familiarity may have contributed to different degrees in the two groups across

10 the conditions so that their estimated familiarity and recollection scores were distorted differently.

11 Although our results suggest that dysfunction in medial temporal lobe structures, perhaps

12 particularly the hippocampus, contributes in a major way to the memory problems in mild to

13 moderate non-dementing PD, this conclusion should be viewed with some caution for several

14 reasons and some qualifications need to be made.

15 First, the idea of compensating for impaired executive functions by providing easy-to-follow

16 instructions remains to be fully explored. It could be that other compensating tasks will be more

17 likely to lead to a disproportionate improvement in PD recall. Even without such disproportionate

18 improvement, however, patients may find that using better but simple encoding and retrieval

19 procedures produces valuable benefits to everyday recall abilities.

20 Second, we are not yet sure which kinds of frontally-related executive function disruptions relate

21 most closely to memory impairments. For example, apathy, a common feature of PD (e.g., Pluck &

22 Brown, 2002; Dujardin, *et al.,* 2007; Barone, *et al.,* 2009), contributes to impaired memory (e.g.,

23 Butterfield, Cimino, Oelke, Hauser & Sanchez-Ramos, 2010) and future studies should be careful to

24 control for this.

1 Third, as previously discussed, there is some evidence that some frontal lesions can impair

2 familiarity. It seems unlikely that this deficit is related to executive function impairment and more

3 likely that the effective damage is to frontal sites that form part of the perirhinal cortex familiarity

4 memory system. However, precisely what the relevant frontal region is and what its exact

5 familiarity-related function is has not yet been explored.

6 Fourth and related to the previous point, future research will need to use structural and functional

7 MRI to identify the extent to which damage or dysfunction of the hippocampus, perirhinal cortex,

8 parts of the frontal lobes, or other structures relate to recollection/recall and familiarity deficits in

9 PD patients. In such research, it will also be critical to measure structure and functionality of the

10 midbrain dopaminergic systems, damage to which underlies PD.

11 Fifth, PD is a heterogeneous disorder. Disruption of familiarity and recollection is likely to be

12 influenced by these variable factors, such as the severity of depression, executive functions, and the

13 severity of different kinds of medial temporal lobe dysfunction. This occurs because PD is a

14 syndrome, defined in terms of its characteristic motor symptoms, the severity of which does not

15 always related straightforwardly to cognitive symptoms that are caused by the atrophy of related

16 but distinct structures. In addition, PD patients are treated with a variety of drugs and there is direct

17 evidence that particular drugs can disrupt recall/recollection (e.g., Edelstyn *et al.,* 2010; MacDonald

18 *et al.,* 2013; Shepherd *et al.,* 2013).

19 Finally, familiarity and recollection are hard to measure so great care must be taken in their

20 measurement. Recollection can be measured directly (as with our source recall measure) and this

21 should be done where possible. But familiarity should either be measured with the RK procedure or

22 a modification of this, such as the familiarity only procedure (see Mayes *et al.,* 2007), ensuring that

23 very careful instructions are given and that checks are made throughout to ensure that participants

24 are following the instructions to the letter.

1 In summary, medicated, non-dementing mild-to-moderate Parkinson's disease (PD) patients

2 exhibited impairments in source recall and subjective recollection as well as a marginal and less

3 severe overall decline in familiarity. Providing full guidance at encoding and retrieval improved

4 source recall and subjective recollection to the same extent in PD and their age matched controls so

5 that PD subjective recollection and source recall remained deficient. On the other hand, familiarity

6 was unaffected by guidance provided at either encoding alone or at retrieval as well as at encoding.

7 The PD pattern of recollection and familiarity response to strategic guidance suggests that their free

8 recall, subjective recollection and source recall impairments are amnesia-like deficits caused at least

9 in part by damage or dysfunction to the hippocampus whereas their milder familiarity impairment

10 may have been caused by perirhinal cortex damage or dysfunction. However, the patients' response

11 to guidance also suggests that their recall and recollection impairments may often arise partly

12 because of a kind of dysexecutive problem caused by damage or dysfunction of parts of the

13 prefrontal cortex. Either way our results provide the hope that poor PD recall and recollection may

14 be usefully helped by given patients guidance to steer more effective encoding and retrieval.

15

16 Acknowledgements and funding

19

35

1 References

2 Aggleton, J.P. & Brown, M.W. (1999). Episodic memory, amnesia, and the hippocampal-anterior
3 thalamic axis. *Behavioural Brain Sciences, 122*, 425-89.

4 Aggleton, J.P., Vann, S.D., Denby, C., Dix, S., Mayes, A.R., Roberts, N., et al. (2005). Sparing of the
5 familiarity component of recognition memory in a patient with hippocampal pathology.
6 *Neuropsychologia*, *43*, 1810–1823.

7 Alexander, G.E., Delong, M.R. & Strick, P.L. (1986). Parallel Organization of Functionally Segregated
8 Circuits Linking Basal Ganglia and Cortex. *Annual Review of Neuroscience*, *9*, 357-381

9 Algarabel, S., Escudero, J., Peset, V., Combita, L-M., Rodriguez, L-A., Fuentes, M., *et al.* (2010).
10 Recognition by familiarity is preserved in Parkinson's without dementia and Lewy-Body disease.
11 *Neuropsychology, 24*, 599-607.

12 Alfonso Recio, L., Martin, P., Carvajal, F., Ruiz, M. & Serrano, J.M. (2013). A holistic analysis of
13 relationships between executive function and memory in Parkinson's disease. *Journal of Clinical and*
14 *Experimental Neuropsychology, 35*, 147-159.

15 Anderson, N.D., Ebert, P.L., Jennings, J.M., Grady, C.L., Cabeza, R. & Graham, S.J. (2008).
16 Recollection- and familiarity-based memory in healthy aging and amnestic mild cognitive
17 impairment. *Neuropsychology, 22*, 177-187.

18 Apostolova, L.G., Beyer, M., Green, A.E., Hwang, K.S., Morra, J.H., Chou, Y.Y., *et al.* (2010).
19 Hippocampal, caudate and ventricular changes in Parkinson's disease with and without dementia.
20 *Movement Disorders*, *25*, 687–688.

21 Barone, P., Antonini, A., Colosimo, C., Marconi, R., Morgante, L., Avarello, T.P., Bottacchi, E., *et al.*
22 (2009) The PRIAMO study: A multicenter assessment of nonmotor symptoms and their impact on
23 quality of life in Parkinson's disease. *Movement Disorders*, *24*, 1641–1649.

24 Bastin, C., Van der Linden, M., Schnakers, C., Montaldi, D. & Mayes, A.R. (2010). The contribution of
25 familiarity to within- and between-domain associative recognition memory: Use of a modified
26 remember/know procedure. *European Journal of Cognitive Psychology*, *22*, 922-943.

27 Baxter, L.R., Schwartz, J.M., Phelps, M.E., Mazziota, J.C., Guze, B.H., Selin, C.E., *et al.* (1989).
28 Reduction of prefrontal cortex glucose metabolism common to three types of depression. *Archives*
29 *of General Psychiatry*, 46, 243-250.

30 Beatty, W.W., Staton, R.D., Weir, W.S., Monson, N. & Whitaker, H.A. (1989). Cognitive disturbances
31 in Parkinson's disease. *Journal of Geriatric Psychiatry and Neurology, 2*, 22-33.

32 Beatty, W.W. & Monson, N. (1990). Problem Solving in Parkinson's Disease: Comparison of
33 Performance on the Wisconsin and California Card Sorting Tests. *Journal of Geriatric Psychiatry and*
34 *Neurology, 3*, 163-71.

Bethus. I., Tse, D. & Morris, R.G. (2010). Dopamine and memory: modulation of the persistence of memory for novel hippocampal NMDA receptor dependent paired associates. *The Journal of Neuroscience*, *30*, 1610 –1618.

Bouchard,T.P., Malykhin, N., Martin, W.R., Hanstock, C.C., Emery, D.J., Fisher, N.J., *et al.* (2008). Age and dementia-associated atrophy predominates in the hippocampal head and amygdala in Parkinson's disease. *Neurobiology of Aging, 29*, 1027–1039.

Bowles, B., Crupi, C., Mirsattari, S.M., Pigott, S.E., Parrent, A.G. & Pruessner, J.C. (2007). Impaired familiarity with preserved recollection after anterior temporal-lobe resection that spares the hippocampus. *Proceedings of the National Academy of Science (USA), 104*, 16382–16387.

Braak, H., Del Tredici, K., Rüb, U., de Vos, R.A.I., Jansen Steur, E.N.H. & Braak E. (2006). Staging of brain pathology related to sporadic Parkinson's disease. *Neurobiology of Aging, 24*, 197–211

Braak, H., Rüb, U. & Del Tredici, K. (2006). Cognitive decline correlates with neuropathological stage in Parkinson's disease. *Journal of Neurological Sciences, 248*, 255–258.

Braak, H. & Del Tredici, K. (2008). Reply to "Controversies over the staging of α-synuclein pathology in Parkinson's disease". *Acta Neuropathology,* 116, 129–131

Breen, E.K. (1993). Recall and recognition memory in Parkinson's disease. *Cortex, 29*, 91-102.

Bruck, A., Kurki, T., Kaasinen, V., Vahlberg, T. & Rinne, J.O. (2004). Hippocampal and prefrontal atrophy with early non-demented Parkinson's disease is related to cognitive impairment. *Journal of Neurology, Neurosurgery and Psychiatry, 75*, 1467-1469.

Bunzeck, N., Schütze, H., Stallforth, S., Kaufmann, J., Düzel, S., Heinze, H-J., *et al.* (2007). Mesolimbic novelty processing in older adults. *Cerebral Cortex, 17*, 2940–2948

Burgess, P.W. & Shallice, T. (1997). *The Hayling and Brixton Tests*. Thames Valley Test Co. Ltd: Bury St. Edmunds.

Butterfield, L., Cimino, C.R., Oelke, L.E., Hauser, R.A. & Sanchez-Ramos, J. (2010). The independent influence of apathy and depression on cognitive functioning in Parkinson's disease. *Neuropsychology*, *24*, 721-730.

Camicioli, R., Moore, M.M., Kinney, A., Corbridge, E., Glassberg, K. & Kaye JA. (2003). Parkinson's disease is associated with hippocampal atrophy. *Movement Disorders, 18*, 784–790.

Carlesimo, G.A., Piras, F., Assogna, F., Pontieri, F.E., Caltagirone, C. & Spalletta, G. (2012). Hippocampal abnormalities and memory deficits in Parkinson's disease. *Neurology*, *78*, 1939-1945.

Channon, S. & Green, P.S.S. (1999). Executive function in depression: the role of performance strategies in aiding depressed and non-depressed participants*. Journal of Neurology, Neurosurgery and Psychiatry, 66*, 162-171.

Chowdhury, R., Guitart-Masip, M., Bunzeck, N., Dolan, R.J. & Duzel, E. (2012). Dopamine modulates episodic memory persistence in old age. *The Journal of Neuroscience, 32*, 14193-14204.

1 Clelland, C.D., Choi, M., Romberg, C., Clemenson, G.D., Fragniere, A., Tyre, P., *et al.* (2009). A
2 functional role for adult hippocampal neurogenesis in spatial pattern separation. *Science*, *325,* 210-
3 213.

4 Cohn, M., McAndrews, M.P. & Moscovitch, M. (2009). Associative reinstatement: A novel approach
5 to assessing associative memory in patients with unilateral temporal lobe excisions.
6 *Neuropsychologia*, *47*, 2989–2994.

7 Cohn, M., Moscovitch, M. & Davidson, P.S.R. (2010). Double dissociation between familiarity and
8 recollection in Parkinson's disease a function of encoding tasks. *Neuropsychologia*, *48*, 4142-4147.

9 Cooper, J.A., Sagar, H.J., Jordan, N., Harvey, N.S. & Sullivan, E.V. (1991). Cognitive impairment in
10 early untreated Parkinson's disease and its relationship to motor disability. *Brain*, *114*, 2095-2122.

11 Crescentini ,C., Mondolo, F., Biasutti, E. & Shallice T. (2008). Supervisory and routine processes in
12 noun and verb generation in nondemented patients with Parkinson's disease. *Neuropsychologia, 46*,
13 434-447.

14 Davidson, P.S.R., Anaki, D., Saint-Cyr, J.A., Chow, T.W. & Moscovitch, M. (2006). Exploring the
15 recognition memory deficit in Parkinson's disease: estimates of recollection versus familiarity. *Brain*,
16 *129*, 1768-1779.

17 Della Rocchetta, A.I. & Milner, B. (1993). Strategic search and retrieval inhibition: The role of the
18 frontal lobes. *Neuropsychologia*, *31*, 503-24.

19 Dirnberger, G. & Jahahshahi, M. (2013). Executive dysfunction in Parkinson's Disease: A review.
20 *Journal of Neuropsychology*, *7*, 193-224.

21 Duarte, A., Ranganath, C. & Knight, R.T. (2005). Effects of unilateral prefrontal lesions on familiarity,
22 recollection and source memory. *The Journal of Neuroscience*, *25*, 8333-37.

23 Dujardin, K., Sockeel, P., Devos, D., Delliaux, M., Krystkowiak, P., Destee, A., *et al.* (2007).
24 Characteristics of apathy in Parkinson's disease. *Movement Disorders, 22*, 778–784.

25 Edelstyn, N.M.J., Mayes, A.R., Condon, L., Tunnicliffe, M. & Ellis, S.J. (2007). Recognition,
26 recollection, familiarity and executive function in patients with moderate Parkinson's Disease.
27 *Journal of Neuropsychology*, *1*, 131-47.

28 Edelstyn, N.M.J., Shepherd, T.A., Mayes, A.R., Sherman, S.M. & Ellis, S.J. (2010). Effect of disease
29 severity and dopaminergic medication on recollection and familiarity in patients with idiopathic
30 nondementing Parkinson's. *Neuropsychologia, 48*, 1367–1375.

31 Edelstyn, N.M.J., Grange, J.A., Ellis, S.J. & Mayes, A.R. (2015). A deficit in familiarity-driven
32 recognition in a right-sided mediodorsal thalamic lesion patient. *Neuropsychology, In press.*

33 Fahn, S. & Elton, R.L. (1987).T he Unified Parkinson's Disease Rating Scale IN: S. Fahn, C.D. Marsden,
34 D.B. Calne, M. Goldstein (Eds.), Recent developments in Parkinson's disease, Macmillan, Florham
35 Park, N.J., pp. 293–304

1   Fearnley, J.M. & Lees, A.J. (1991). Ageing and Parkinsons disease: Substantia nigra regional
2   selectivity. *Brain*, *114*, 2283-2301.

3   Filoteo, J.V., Reed, J.D., Litvan, I. & Harrington, D.L. (2014). Volumetric correlates of cognitive
4   functioning in nondemented patients with Parkinson's disease. *Movement Disorders, 29*, 361-367.

5   Flowers, K.A., Pearce, I. & Pearce, J.M.S. (1984). Recognition memory in Parkinson's disease. *Journal*
6   *of Neurology, Neurosurgery and Psychiatry*, *47*, 1174-81.

7   Folstein, M.F., Folstein, S.E. & McHugh, P.R. (1975). Mini-mental state: A practical method for
8   grading the cognitive state of patients for the clinician. *Journal of Psychiatry Research, 12*, 189-98.

9   Gabrieli, J.D.E., Singh, J., Stebbins, G.T. & Goetz, C.G. (1996). Reduced working memory span in
10   Parkinson's disease: Evidence for the role of a frontostriatal system in working and strategic
11   memory. *Neuropsychology*, *10*, 322-32.

12   Gasbarri, A., Sulli, A. & Packard, M.G. (1997). The dopaminergic mesencephalic projections to the
13   hippocampal formation in the rat. *Progress in Neuro-Psychopharmacology and Biological Psychiatry*,
14   *21*, 1-22.

15   Gershberg, F.B. & Shimamura, A.P. (1995). Impaired use of organizational strategies in free recall
16   following frontal lobe damage. *Neuropsychologia, 33*, 1305-33.

17   Gradin, V.B. & Pom, A. (2008). The role of hippocampal atrophy in neuropsychiatric disorders.
18   *Journal of Biological Physics, 34,* 107-120.

19   Hammad, H. & Wagner, J.J. (2006). Dopamine-mediated disinhibition in the CA1 region of rat
20   hippocampus via D_3 receptor activation. *The Journal of Pharmacology and Experimental*
21   *Therapeutics, 316,* 113-120.

22   Hanes, K.R., Andrewes, D.G., Smith, D.J. & Pantelis, C. (1996). A brief assessment of executive control
23   dysfunction: Discriminant validity and homogeneity of planning, set shift, and fluency measures.
24   *Archives of Clinical Neuropsychology, 11*, 185-91.

25   Harlow, I.M., MacKenzie G. & Donaldson, D.A. (2010). Familiarity for associations? A test of the
26   domain dichotomy theory. *Journal of Experimental Psychology: Learning, Memory and Cognition*, *36,*
27   1381-1388.

28   Hay, J.F., Moscovitch, M. & Levine, B. (2002). Dissociating habit and recollection: evidence from
29   Parkinson's disease, amnesia and focal lesion patients. *Neuropsychologia*, *40,* 1324-34.

30   Henson, R.N.A., Rugg, M.D., Shallice, T., Josephs, O. & Dolan, R.J. (1999). Recollection and familiarity
31   in recognition memory: An event-related functional magnetic resonance imaging study. *The Journal*
32   *of Neuroscience, 19*, 3962-3972.

33   Higginson, C.I., Wheelock, V.L., Carroll, K.E. & Sigvardt, K.A. (2005). Recognition memory in
34   Parkinson's disease with and without dementia: Evidence inconsistent with the retrieval deficit
35   hypothesis. *Journal of Clinical and Experimental Neuropsychology*, *27*, 516-28.

1 Hirst, W. & Volpe, B.T. (1988). Memory strategies with brain damage. *Brain and Cognition*, *8*, 379-
2 408.

3 Hoehn, M.M. & Yahr, M.D. (1967). Parkinsonism: Onset, progression and mortality. *Neurology*, *17*,
4 427–42.

5 Holdstock, J.S., Mayes, A.R., Roberts, N., Cezayirli, E., Isaac, C.L. & O'Reilly, R.C. (2002). Under what
6 conditions is recognition spared relative to recall after selective hippocampal damage in humans.
7 *Hippocampus*, *12*, 341–351.

8 Hornykiewicz, O. (1966). Dopamine (3-hydroxytyramine) and brain function. *Pharmacology Review,*
9 *18*, 925-64.

10 Ibarretxe-Bilbao, N., Ramirez-Ruiz, B., Tolosa, E., Marti, M.J., Valldeoriola, F., Bargallo, N., *et al.*
11 (2008). Hippocampal head atrophy predominance in Parkinson's disease with hallucinations and
12 with dementia. *Journal of Neurology, 255*, 1324–1331.

13 Jacoby, L.L. (1991). A process dissociation framework: separating automatic from intentional uses of
14 memory. *Journal of Memory and Language, 30*, 513-41.

15 Janowsky, J.S., Shimamura, A.P. & Squire, L.R. (1989a). Source memory impairment in patients with
16 frontal lobe lesions. *Neuropsychologia, 27*, 1043-56.

17 Janowsky, J.S., Shimamura, A.P., Kritchevsky, M. & Squire, L.R. (1989b). Cognitive impairment
18 following frontal lobe damage and its relevance to human amnesia. *Behavioural Neuroscience, 103*,
19 548-60.

20 Jetter, W., Poser, U., Freeman, R.B. Jr & Markowitsch, H.J. (1986). A verbal long-term memory deficit
21 in frontal lobe damaged patients. *Cortex*, *22*, 229–242.

22 Junqué, C., Ramīrex-Ruiz, B., Tolosa, E., Summerfield, C., Marti, M.J., Pastor, P., *et al.* (2005).
23 Amygdalar and hippocampal MRI volumetric reductions in Parkinson's disease with dementia.
24 *Movement Disorders, 20*, 540–544.

25 Kaftas, A. & Montaldi, D. (2014). Two separate, but interacting, neural systems for familiarity and
26 novelty detection: A dual-route mechanism. *Hippocampus, 24*, 516-527.

27 Kemperman, G. & Kroneberg, G. (2003). Depressed new Neurons?—Adult hippocampal
28 neurogenesis and a cellular plasticity hypothesis of major depression. *Biological Psychiatry*, *54*, 499-
29 503.

30 Knoke, D., Taylor, A.E. & Saint-Cyr, J.A. (1998). The differential effects of cueing on recall in
31 Parkinson's disease and normal subjects. *Brain and Cognition, 38*, 261-274.

32 Laakso, M.P., Partanen, K., Riekkinen, P., Lehtovirta, M., Helkala, E.L., Hallikainen, M., *et al.* (1996).
33 Hippocampal volumes in Alzheimer's disease, Parkinson's disease with and without dementia, and in
34 vascular dementia: an MRI study. *Neurology*, *46*, 678-81.

35 Laszy, J., Laszlovszky, I. & Gyertyan, I. (2005). Dopamine D_3 receptor antagonists improve the
36 learning performance in memory-impaired rats. *Psychopharmacology*, *179*, 567-575.

1 Lisman, J.E. & Grace, A.A. (2005). The hippocampal-VTA loop: controlling the entry of information
2 into long-term memory. *Neuron*, *46*, 703–713.

3 Lisman, J.E., Grace, A.A. & Duzel, E. (2011). A neoHebbian framework for episodic memory; role of
4 dopamine-dependent late LTP. *Trends in Neuroscience, 34,* 536 –547.

5 MacDonald, A.A., Seergobin, K.N., Owen, A.M., Tamjeedi, R., Monchi, O., Ganjavi, H., *et al.* (2013).
6 Differential effects of Parkinson's disease and dopamine replacement on memory encoding and
7 retrieval. *Plos One*, *8*, 1-9.

8 MacPherson, S.E., Bozzali, M., Cipolotti, L., Dolan, R.J., Rees, J.H. & Shallice, T. (2008). Effect of
9 frontal lobe lesions on the recollection and familiarity components of recognition memory.
10 *Neuropsychologia*, *46*, 3124-32.

11 Mayes, A.R. (1988). Human Organic Memory Disorders. Cambridge University Press.

12 Mayes, A.R. & Daum, I. (1996). How specific are the memory and other cognitive deficits caused by
13 frontal lobe lesions? P. Rabbitt, Editor, Methodology of Frontal and Executive Functions. Hove:
14 Erlbaum.

15 Mayes, A.R., Meudell, P.R. & Neary, D. (1980). Do amnesics adopt inefficient encoding strategies
16 with faces and random shapes? *Neuropsychologia*, *18*, 527-541.

17 Mayes, A.R., Montaldi, D. & Migo, E.M. (2007). Associative memory and the medial temporal lobes.
18 *Trends in Cognitive Sciences, 11*, 126–135

19 Migo, E.M., Mayes, A.R. & Montaldi, D. (2012). Measuring recollection and familiarity: Improving
20 the remember/know procedure. *Consciousness and Cognition, 21*, 1435-1455.

21 Montaldi, D., Spencer, T.J., Roberts, N. & Mayes, A.R. (2006). The neural system that mediates
22 familiarity memory. *Hippocampus, 16*, 504-520.

23 Montaldi, D. & Mayes, A.R. (2010). The role of recollection and familiarity in the functional
24 differentiation of the medial temporal lobes. *Hippocampus*, *20*, 1291-1314.

25 Morris, R.G., Downes, J.J., Sahakian, B.J., Evenden, J.L., Heald, A. & Robbins, T.W. (1988). Planning
26 and spatial working memory in Parkinson's disease. *Journal of Neurology, Neurosurgery and
27 Psychiatry, 51*, 757-66

28 Nagano-Saito, A., Washimi, Y., Arahata, Y., Kachi, T., Lerch, J.P., Evans, A.C., *et al.* (2005). Cerebral
29 atrophy and its relation to cognitive impairment in Parkinson's disease. *Neurology, 64*, 224-229.

30 O'Carroll, C.M., Martin, S.J., Sandin, J., Frenguelli, B. & Morris, R.G. (2006). Dopaminergic modulation
31 of the persistence of one-trial hippocampus-dependent memory. *Learning and Memory, 13*, 760–
32 769

33 Obeso, J.A., Cruz Rodriguez-Oroz, M., Benitez-Temino, B., Blesa, F.J., Guridi, J., Marin, C., *et al.*
34 (2008). Functional organisation of the basal ganglia: Therapeutic implications for Parkinson's
35 disease. *Movement Disorders*, *23*, S548-S559.

1 Owen, A.M., Doyon, J., Dagher, A., Sadikot, A. & Evans, A.C. (1998). Abnormal basal ganglia outflow
2 in Parkinson's disease identified with PET. Implications for higher cortical functions. *Brain*, *121*, 949-
3 965.

4 Parkin, A.J., Bindschaedler, C., Harsent, L. & Metzler, C. (1996). Pathological false alarm rates
5 following damage to the left frontal cortex. *Brain and Cognition*, *32* 14-27

6 Parkinson, J. (2002). An Essay on the Shaking Palsy. *Journal of Neuropsychiatry and Clinical*
7 *Neurosciences, 14*, 223-236.

8 Pereria, J.B., Junqué, C., Bartrés-Faz, D., Ramīrex-Ruiz, B., *et al.* (2013). Regional vulnerability of
9 hippocampal subfields and memory deficits in Parkinson's disease. *Hippocampus, 23*, 720-728.

10 Pluck, G.C. & Brown, J. (2002). Apathy in Parkinson's disease. *Neurology, Neurosurgery and*
11 *Psychiatry, 73*, 636-642.

12 Reynolds, W.M. & Koback, K.A. (1995). Reliability and validity of the Hamilton Depression Inventory:
13 A paper-and-pencil version of the Hamilton Depression Rating Scale Clinical Interview. Psychological
14 Assessment.

15 Rodríguez, R.A., Algarabel, S. & Escudero, J. (2014). Exploring recollection and familiarity
16 impairments in Parkinson's disease. *Journal of Clinical and Experimental Neuropsychology, 36*, 494-
17 506.

18 Riekkinen, P., Kejonen, K., Laakso, M.P., Soininen, H., Partanen, K. & Riekkinen, M.   (1998).
19 Hippocampal atrophy is related to impaired memory, but not frontal functions in non-demented
20 Parkinson's disease patients. *NeuroReport*, *9*, 1507–1511.

21 Sadeh, T., Ozubko, J.D., Winocur, G. & Moscovitch, M. (2014). How we forget may depend on how
22 we remember. *Trends in Cognitive Science, 18*, 26-36.

23 Sahay, A. & Hen, R. (2007). Adult hippocampal neurogenesis in depression. *Nature Neuroscience, 10*,
24 1110-1115.

25 Sahay, A., Scobie, K.N., Hill, A.S., O'Carroll, C.M., Kheirbek, M.A., Burghardt, N.S., et al. (2011).
26 Increasing adult hippocampal neurogenesis is sufficient to improve pattern separation. *Nature*, *472*, 466-470.

27 Saksida, L.M. & Bussey, T.J. (2010). The representational-hierarchical view of amnesia: translation
28 from animal to human. *Neuropsychologia, 48*, 2370–2384.

29 Savage, C.R., Deckersbach, T., Heckers, S., Wagner, A.D., Schacter, D.L., Alpert, N.L., *et al*. (2001).
30 Prefrontal regions supporting spontaneous and directed application of verbal learning strategies.
31 Evidence from PET. *Brain, 124,* 219-231.

32 Schacter, D.L., Curran, C., Galluccio, L., Milberg, W.P. & Bates, J.F. (1996). False recognition and the
33 right frontal lobe: A case study. *Neuropsychologia, 34,* 793-808.

1 Segura, B., Ibarretxe-Bilbao, N., Sala-Llonch, R., Baggio, H.C., Marti, M.J., Valldeoriola, F., *et al.*
2 (2012). Progressive changes in a recognition memory network in Parkinson's disease. *Journal of*
3 *Neurology, Neurosurgery and Psychiatry*, doi: 10.1136/jnnp-2012-302822

4 Shepherd, T.A., Edelstyn, N.M.J., Mayes, A.R. & Ellis, S.J. (2013). Second Generation Dopamine
5 Agonists and Recollection Impairments in Parkinson's disease. *Journal of Neuropsychology*, *7,* 284-
6 305*.*

7 Shimamura, A.P., Jurica, P.J., Mangels, J.A. & Gershberg, F.B. (1995).Susceptibility to memory
8 interference effects following frontal lobe damage: findings from tests of paired-associate learning.
9 *Journal of Cognitive Neuroscience,* 7, 144-52.

10 Snodgrass, J.G. & Corwin, J. (1988). Pragmatics of measuring recognition memory. Applications to
11 dementia and amnesia. *Quarterly Journal of Experimental Psychology A*, *117*, 34-50.

12 Tam, C.W., Burton, E.J., McKeith, I.G., Burn, D.J. O'Brien, J.T. (2005). Temporal lobe atrophy on MRI
13 in Parkinson disease with dementia: a comparison with Alzheimer disease and dementia with Lewy
14 bodies. *Neurology*, *64*, 861–865.

15 Takahashi, H., Kato, M., Takano, H., Arakawa, R., Okumura, M., Otsuka, T., *et al*. (2008). Differential
16 contributions of prefrontal and hippocampal dopamine D1 and D2 receptors in human cognitive
17 functions. *The Journal of Neuroscience, 28*, 12032-12038.

18 Taylor, A.E., Saint-Cyr, J.A. & Lang, A.E. (1986). Frontal lobe dysfunction in Parkinson's disease and
19 depression. *Brain*, *109*, 845-883.

20 Tulving, E. (1985). Memory and consciousness. *Canadian Psychologist*, *26*, 1-12.

21 Turner, M.S., Cipolotti, L., Yousry, T.A. & Shallice, T. (2007). Confabulation: damage to a specific
22 inferior medial prefrontal system. *Cortex*, *44*, 637-48.

23 Van Spaendonck, K.P.M., Berger, H.J.C., Horstink, M.W.I.M., Borm, G.F. & Cools, A.R. (1996).
24 Memory performance under varying cueing conditions in patients with Parkinson's disease.
25 *Neuropsychologia, 34*, 1159-1164.

26 Wechsler, D. (1997). *Wechsler Memory Scale III.* The Psychological Corporation, Harcourt Brace & Co.

27 Wechsler, D. (1999). *Wechsler Abbreviated Scale of Intelligence*. The Psychological Corporation,
28 Harcourt Brace & Co.

29 Weiermann, B., Stephan, M.A., Kaelin-Lang, A. & Meier, B. (2010). Is there a recognition memory
30 deficit in Parkinson's disease: Evidence from estimates of recollection and familiarity. *International*
31 *Journal of Neuroscience,* 120, 211-16.

32 Wheeler, M.A., Stuss, D.T. & Tulving, E. (1995). Frontal lobe damage produces episodic memory
33 impairment. *Journal of International Neuropsychological Society*, 1, 525-36.

34 Whittington, C.J., Podd, J. & Kan, M.M. (2000). Recognition memory impairment in Parkinson's
35 disease: Power and meta-analyses. *Neuropsychology*, 14, 233-46.

1   Wixted, J.T. & Squire, L.R. (2011). The medial temporal lobes and attributes of memory. *Trends in*
2   *Cognitive Science,* 15, 210-217.

3   Yonelinas, A.P. (2002). The nature of recollection and familiarity: A Review of 30 Years of Research.
4   *Journal of Memory and Language,* 46, 441-517.

5   Yonelinas, A.P. & Jacoby, L.L. (1995). The relation between remembering and knowing as bases for
6   recognition: Effects of size congruency. *Journal of Memory and Language, 34*, 622-43.

Table 1. The demographic, neuropsychological and clinical (patients only) characteristics for the groups of controls and patients.

Parameter	Healthy controls (n=22)	Parkinson's patients (n=30)
	Mean (SD)	Mean (SD)
Age (years)	65.50 (5.25)	64.38 (6.51)
Current levels of functioning		
MMSE[1]	29.00 (0.95)	29.06 (1.12)
WASI[2] (full scale IQ)	108.83 (8.67)	109.69 (14.1)
Illness duration (years)	----	6.31 (3.34)
Modified Hohn and Yahr disease severity rating	----	2.53 (0.90)
UPDRS[3] (motor subsection)	----	13.38 (5.08)
Equivalent Dopamine Load (mg/day)	----	635.71 (463.86)

Notes and abbreviations: [1]MMSE, Mini Mental State Examination; [2]WASI, Wechsler Abbreviated Scales of Intelligence; [3]UPDRS, motor subsection of the Unified Parkinson's Disease Rating Scale.

Table 2. Mean hit rate, false alarm rate for recognition memory, know, remember, source recall, and estimates of recognition memory (d'), familiarity (d'), subjective recollection (pr) and source recall (pr) are shown for the groups of controls and patients for each the three experimental recognition memory test conditions.

|	Recognition memory			Know			Remember			Source recall		
	HR	FAR	RM (d')	HR	FAR	Fam (d')	HR	FAR	Recoll (pr)	HR	FAR	Source (pr)
Condition 1: Spontaneous encoding and retrieval												
Controls	37.41	4.23	2.23	10.45	2.55	2.02	25.64	1.86	0.47	19.41	2.41	0.30
1 SD	7.34	2.71	0.89	4.55	2.20	2.14	9.52	2.03	0.22	5.42	2.12	0.10
Patients	28.67	5.17	1.55	17.20	4.70	1.25	10.50	0.57	0.20	8.47	0.90	0.15
1 SD	7.98	4.03	0.53	8.00	4.12	0.53	634.00	0.94	0.12	7.34	1.49	0.15
Condition 2: Partial guidance (encoding only)												
Controls	36.95	3.73	2.33	9.00	2.64	1.51	24.32	1.95	0.5	20.56	2.88	0.33
1 SD	7.85	2.71	0.63	3.99	2.50	0.62	10.83	1.76	0.22	6.75	2.91	0.17
Patients	30.43	3.50	1.87	11.77	3.30	1.23	15.57	0.40	0.30	12.87	0.33	0.25
1 SD	8.85	3.36	0.58	6.12	3.57	0.69	6.35	0.77	0.13	8.92	0.66	0.18
Condition 3: Full guidance (encoding and retrieval)												
Controls	37.75	2.81	2.38	6.31	2.13	1.34	30.38	1.50	0.57	23.00	2.09	0.41
1 SD	8.39	1.83	0.53	4.21	1.63	0.53	11.47	0.68	0.23	9.00	2.07	0.18
Patients	28.7	2.80	1.90	9.83	2.47	1.22	16.93	0.33	0.33	13.45	0.43	0.26
1 SD	7.72	5.57	0.56	4.71	3.42	0.42	6.63	0.66	0.13	9.37	0.63	0.19

Notes and abbreviations: HR, FAR, Hit rate and False alarm rate, respectively; Recoll, subjective recollection; Source, source recall; d', signal detection mesure of discrimination accuracy; pr, threshold measure of accuracy; 1 SD, one standard deviation. Six PD patients and four controls were lost to follow up between conditions 2 and 3.

Table 3. Neuropsychological and depression scores for the groups of controls and patients.

Parameter	Healthy controls (n=17)		Parkinson's patients (n=10)
	Mean (SD)		Mean (SD)
Brixton Test	5.85 (0.8)		3.39 (2.51)***
Logical Memory (30 min delay)	31.39 (4.48)		21.59 (7.05)***
Hamilton Depression Inventory	4.61 (2.93)		11.92 (6.16)**

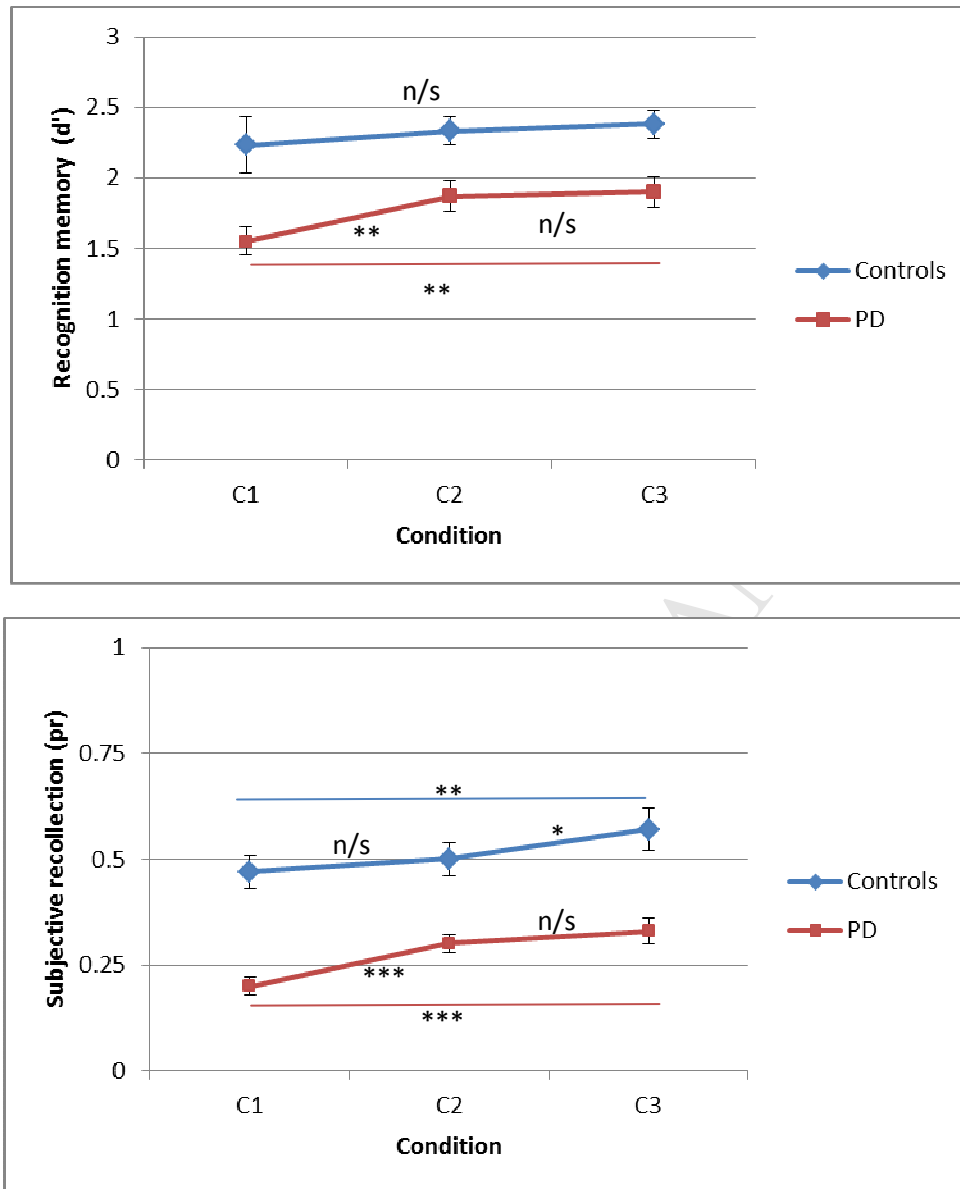Notes and abbreviations: significant at *p<.05, **p<.001, ***p<.001.

Figure 1. Estimates of recognition memory (d'), familiarity (d'), subjective recollection (pr) and source recall (pr) are shown for the groups of controls and patients for the three experimental recognition memory test conditions.
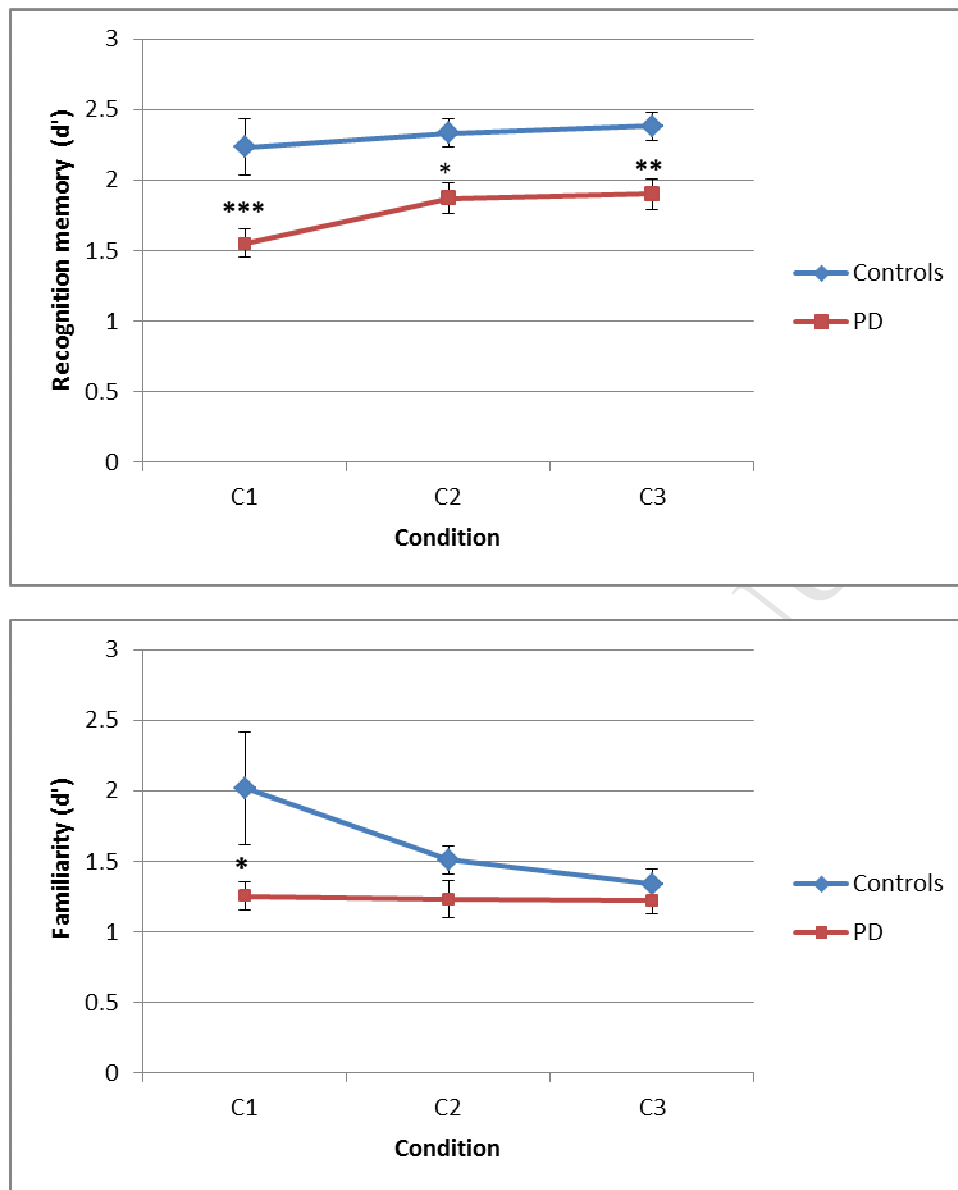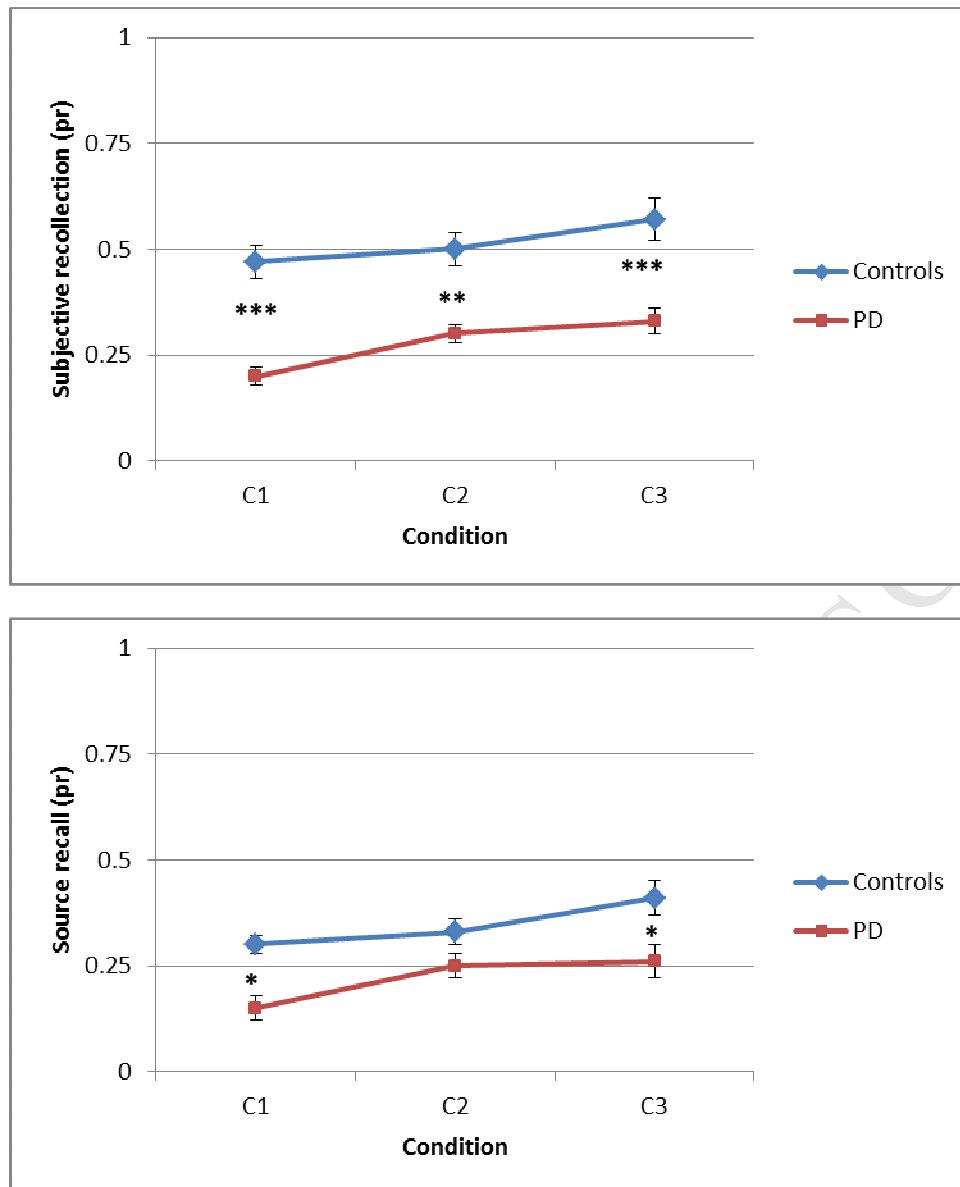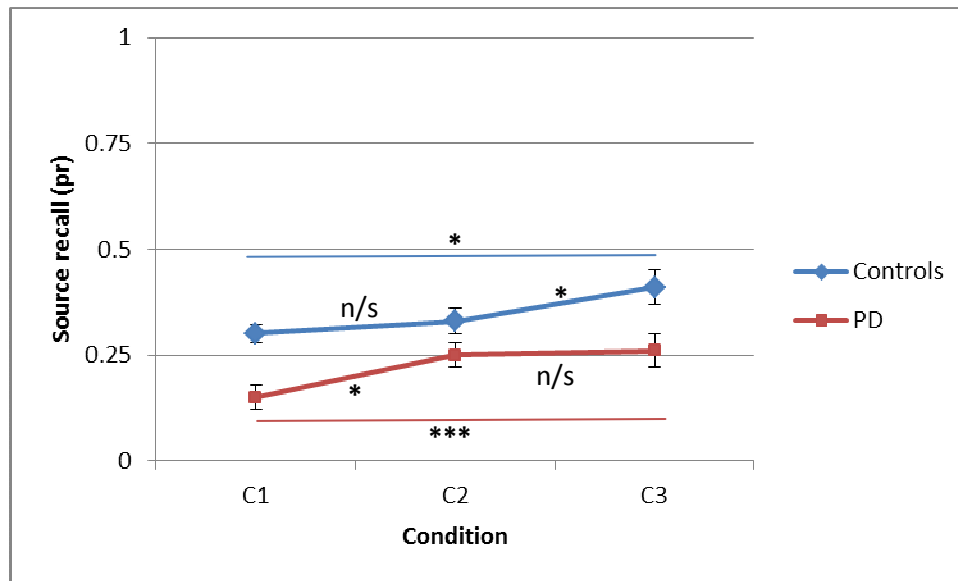
Notes and abbreviations: Significant at *p<.05, **p<.01, ***p<.001. C1, baseline condition where participants are reliant on spontaneously generated encoding and retrieval strategies; C2 and C3 provide guidance at encoding (C2) and additionally at retrieval (C3).

Figure 2. Effect of the three experimental recognition memory test conditions on within group estimates of recognition memory (d'), subjective recollection (pr) and source recall (pr) for the groups of controls and patients.

Notes and abbreviations: Significant at *p<.05, **p<.01, ***p<.001. C1, baseline condition where participants are reliant on spontaneously generated encoding and retrieval strategies; C2 and C3 provide guidance at encoding (C2) and additionally at retrieval (C3). N/s, not significant p > .05.