

Prospective memory function in mild cognitive impairment and early dementia

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Abstract

When compared with controls, both mild cognitive impairment (MCI) and dementia are each associated with impaired memory for future intentions, or prospective memory (PM). However, prior studies have failed to agree on whether there are group differences in PM function between those with MCI and dementia. Furthermore, the degree and nature of the impairment remains to be clarified, as does the degree to which this impairment is secondary to deficits in other aspects of cognition. In the present study, MCI ($n = 48$), dementia ($n = 39$), and control participants ($n = 53$) were compared on *Virtual Week*, a measure that closely represents the types of PM tasks that occur in everyday life. Both clinical groups exhibited impairment irrespective of the specific task demands, but the magnitude of this deficit was greater for those with dementia. After covarying for other key cognitive parameters, although the absolute magnitude of the deficit was reduced, significant impairment remained. These results indicate that individuals with MCI, and to a greater extent dementia, experience generalized difficulties with PM. It is suggested that, while other cognitive deficits contribute to these difficulties, there is something unique to prospective remembering that may be additionally disrupted in these groups. (*JINS*, 2010, 16, 318–325.)

Keywords: Virtual week, Assessment, Memory loss, MCI, Dementia, Alzheimer's disease, Aged

INTRODUCTION

Prospective memory (PM) refers to memory for future intentions, such as remembering to take medication or turn off appliances. This ability is crucial to the maintenance of functional independence, which is a fundamental concern for older adults (Chasteen, Park, & Schwarz, 2001). Problems with PM cause more deficits in activities of daily living, instrumental activities of daily living, and caregiver burden than do retrospective memory failures (Smith, Della Sala, Logie, & Maylor, 2000). It is, therefore, of considerable concern that PM is often disrupted in the context of normal adult aging (Henry, Macleod, Phillips, & Crawford, 2004) and to an even greater extent in dementia, even in the mild (Martins & Damasceno, 2008) and preclinical (Duchek, Balota, & Cortese, 2006; Jones, Livner, & Backman, 2006) stages.

To date, seven dementia studies have examined PM performance and have included controls, thus allowing the presence and magnitude of PM impairment to be quantified. All seven of these studies reported that individuals with dementia exhibit PM difficulties relative to controls (Blanco-Campal, Coen, Lawlor, Walsh, & Burke, 2009; Duchek et al., 2006; Jones et al., 2006; Kazui et al., 2005; Kinsella, Ong, Storey, Wallace, & Hester, 2007; Martins & Damasceno, 2008; Troyer & Murphy, 2007). Particularly striking were the findings of Kinsella et al. (2007). In their study, a simple event-based PM task was administered, in which participants were required to remember to make a word substitution whenever a target word appeared in a passage of text. Despite minimal retrospective memory demands (participants were required to recall only one target word), and a relatively mild level of dementia, the Alzheimer's disease (AD) group performed close to floor-level on this task. Therefore, PM difficulties appear to be a prominent and consistent feature of dementia. Such findings are unsurprising given that the neural structures affected in the most common types of dementia,

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even in the early stages, are also known to be implicated in PM function. In particular, prominent atrophy and tau deposition is observed in temporal and frontal neocortices (e.g., Barnes, Ourselin, & Fox, 2009; Scheltens, 2009). There is evidence that the frontal lobes play a key role in various cognitive control operations, such as planning and monitoring, each of which are considered essential to PM performance (see Glisky, 1996; Reynolds, West, & Braver, 2009; Zöllig, West, Martin, Altgassen, Lemke, & Kliegel, 2007). Furthermore, recent evidence has highlighted the importance of temporal neural structures in subserving many important aspects of PM function (den Ouden, Frith, Frith, & Blakemore, 2005; Reynolds et al., 2009).

Research attention has increasingly focused on clarifying whether PM difficulties may have diagnostic significance for early detection of dementia over and above the contribution of retrospective memory, with most research to date supportive of this possibility. Thus, it has been shown that measures of PM make an independent contribution beyond that of retrospective memory to the diagnosis of dementia (Duchek et al., 2006) and to the prediction of dementia 3 years later (Jones et al., 2006). Specifically, Jones et al. (2006) administered measures of prospective and retrospective memory to preclinical AD and control participants, 3 years before dementia diagnosis. The results indicated the preclinical AD participants were impaired on both measures, and within the PM task itself, the prospective and retrospective components were comparably impaired. Furthermore, the PM task contributed significant unique variance to the prediction of AD. These data were particularly striking given that a relatively insensitive measure of PM was used: a single trial that involved reminding the experimenter to make a telephone call after all the tests had been completed. Duchek et al. (2006) also found that an event-based PM task contributed additional unique variance to discriminating mild dementia from controls, above and beyond measures of retrospective memory.

Another way of addressing how early in the disease process PM difficulties arise, and whether the presence of PM difficulties has diagnostic significance, is to examine those diagnosed with mild cognitive impairment (MCI). MCI is characterized by subjective and objective cognitive decline greater than expected for an individual's age and education level, but which does not cause significant functional impairment (Petersen, 2007). Although there is ongoing debate as to whether MCI represents a prodrome of dementia, relative to the general older adult population, this group has a substantially elevated risk of developing dementia and presents with cognitive and brain changes that are generally intermediate between individuals with dementia and nonclinical controls (Albert & Blacker, 2006; Petersen, 2007).

As in dementia, PM is disrupted in MCI and not simply because of problems with the retrospective component of the PM task (Blanco-Campal et al., 2009; Karantzoulis, Troyer, & Rich, 2009; Kazui et al., 2005; Schmitter-Edgecombe, Woo, & Greeley, 2009; Troyer & Murphy, 2007). Karantzoulis et al. (2009) found that individuals with amnesic-MCI were

impaired relative to controls on measures of time- and event-based PM, and these difficulties reflected failures in both the prospective and retrospective components of the tasks. Schmitter-Edgecombe et al. (2009) found that amnesic and non-amnesic MCI participants were impaired on a simple event-based measure of PM (remembering to request a medicine bottle every time a specific task was completed). Because all participants were able to recall the PM task instructions, the PM failure could not be attributed to problems with the retrospective component of the task. It was also argued that the level of MCI-related impairment on the PM measure was greater than the corresponding impairment observed on a separate measure of retrospective memory. Finally, Blanco-Campal et al. (2009) found that an event-based PM task was superior to two retrospective memory tasks in discriminating between MCI of suspected AD etiology and normal controls.

However, a key issue in clinical practice is whether there are group differences, not only between normal aging and pathology, but between different clinical states. Only two PM studies to date have simultaneously assessed both MCI and dementia in comparison to healthy controls, and these came to different conclusions (Kazui et al., 2005; Troyer & Murphy, 2007). Kazui et al. (2005) compared individuals with MCI, AD and demographically matched controls on the Rivermead Behavioral Memory Test (RBMT; Wilson et al., 1989), which includes three PM subcomponents, each consisting of a single task. Although both the MCI and the dementia groups exhibited PM impairment relative to controls, PM performance was comparable for the two clinical groups. Troyer and Murphy (2007) also found that two PM measures (time- and event-based PM) were impaired in both clinical groups, but those with AD were more impaired than the amnesic MCI group.

It, therefore, remains unclear whether measures of PM are sensitive to group differences between these two clinical conditions. The difference in Mini-Mental State Examination (MMSE) scores between the MCI and AD groups was greater in the study by Kazui et al. (2005) (26.7 *vs.* 21.9) than in the study by Troyer and Murphy (2007) (27.8 *vs.* 25.5), thus the absence of PM performance differences in the former study did not simply reflect more closely overlapping clinical groups. One possibility is that the differences between these studies reflect method variance, and specifically, differences in the sensitivity of the PM measure used to index this construct. McDaniel and Einstein (2007) concluded that most PM tasks lack reliability, with some tasks as low as 20%. It was argued that this lack of reliability is attributable to the few opportunities typically given to perform the PM task, with this in particular an issue for many clinical assessments such as the RBMT. In the study by Kazui et al. (2005), the RBMT was used to index PM, with scores on each one of the three RBMT PM tasks reported separately. By contrast, Troyer & Murphy's (2007) study would have had greater sensitivity, with scores based on a total of eight targets assessed.

The present study's first aim was to assess whether a measure of PM with documented reliability and sensitivity

shows group differences in performance between normal aging, MCI and dementia. We used an adapted version of a board game to test PM, Virtual Week (see below), which has documented sensitivity to age-related cognitive impairment (Rendell & Henry, 2009; Will, Rendell, Ozgis, Pierson, Ong, & Henry, 2008). An important advantage of Virtual Week is the inclusion of PM tasks that vary in their relative task demands. In the context of clinical practice, a differentiated profile of impairment on Virtual Week may be informative of the degree of PM impairment *per se* and, also of the particular circumstances in which PM impairment is more likely to arise (and consequently the manner in which rehabilitation efforts should be targeted). Of the MCI and dementia studies conducted to date, only three have compared performance across multiple task parameters, and consequently helped clarify whether specific types of PM processing are particularly disrupted. Troyer and Murphy (2007) found that, although participants with dementia were equivalently impaired on measures of time- and event-based PM, participants with MCI were particularly impaired on the former. Similarly, while Karantzoulis et al. (2009) found that MCI participants were impaired on both time- and event-based PM, the magnitude of the deficit for the former was nearly twice as large. Finally, Blanco-Campal et al. (2009) manipulated the specificity of the instructions and perceptual salience of the PM cue and found that the nonspecific, non-salient condition was associated with greater MCI-related impairment. Kazui et al. (2005) also presented performance separately across the three individual PM tasks of the RBMT, but did not present any statistical comparison of these task-types. Taken together, these studies suggest that PM tasks that impose relatively greater demands on self-initiated retrieval processes, or strategic resources, may be particularly sensitive to the presence of MCI, although they may not differ in their relative sensitivity to dementia.

The second aim was to further assess whether the specific demands of the PM task interact with group status. Virtual Week not only differentiates between event- and time-based PM, but between regular and irregular tasks. This was the one task distinction that was found by Rendell and Craik (2000) to interact with age, with age-related deficits substantially attenuated on regular compared with irregular tasks. Kliegel, Martin, McDaniel, and Einstein (2002) argue that remembering the content of the many different irregular tasks requires more processing resources than remembering the content of the same two regular tasks each day, and in particular, imposes greater demands on retrospective memory. Consequently, it seems likely that irregular tasks will be more sensitive to the presence of MCI than regular tasks.

Our third aim was to provide an assessment of whether PM as indexed by Virtual Week is sensitive to differences between healthy controls, MCI, and dementia, even after taking into account group differences in retrospective memory, working memory and executive functioning. This assessment was considered important because each of these cognitive abilities are related to PM function in normal adult aging (Martin, Kliegel, & McDaniel, 2003; McDaniel & Einstein,

1992), with preliminary evidence supporting such a relationship in the context of abnormal adult aging (Schmitter-Edgecombe et al., 2009; Troyer & Murphy, 2007; although see Martins & Damasceno, 2008).

METHODS

Participants

Of 140 community dwelling Sydney participants, 39 met Diagnostic and Statistical Manual of Mental Disorder (DSM-IV) criteria for dementia (diagnoses of subtype were not known in all cases but included 10 cases of AD, 2 mixed AD and vascular dementia, 1 Lewy body dementia, and 1 dementia pugilistica. These diagnostic groupings are not explored in this study as the groups would be too small for meaningful analysis), 48 met the modified criteria for MCI (14 amnesic single-domain, 6 amnesic multi-domain, and 28 non-amnesic cases; Artero, Petersen, Touchon, & Ritchie, 2006; Petersen, 2007), and 53 were controls without cognitive impairment. All participants had adequate eyesight, hearing and English language ability for the assessment. Participants were excluded if they had a previous diagnosis of psychiatric or neurological illness. Typical medications included cholinesterase inhibitors (taken by 10 participants in the dementia group), and treatments for physical illnesses. The groups did not differ in age, $F(2,135) = 1.68$; $p = .191$; $\eta_p^2 = .02$, education, $F(2,98) = 0.50$; $p = .611$; $\eta_p^2 = .01$, or gender, $\chi^2(2, N = 140) = 1.77$; $p = .412$; $\phi = .11$), but MMSE scores differentiated the groups, ($F(2,134) = 19.66$; $p < .001$, $\eta_p^2 = .23$; see Table 1). Follow-up Tukey tests indicated that participants with dementia had lower scores on the MMSE relative to controls ($p < .001$), and to the MCI group ($p = .001$), but the MCI and control groups did not differ ($p = .329$). MMSE scores indicate that dementia participants were in the very mild stage of illness, consistent with the majority of this group being less than 2 years after diagnosis.

The majority of participants (> 90%) were recruited from a large epidemiological study of aging (Kochan et al., 2009), which commenced 2 years before the current study. The second source of recruitment was from a Memory Disorders Clinic, with many of these participants newly diagnosed with dementia at the time of recruitment. MCI and dementia were diagnosed by consensus conference of the memory clinic or epidemiological study. One of the authors (HB) was a consultant psychiatrist in both of these conferences, the same neuropsychologist oversaw both, and the test batteries used were identical. The same criteria were applied in both situations.

Procedure

Ethics approval was obtained from the South-Eastern Sydney Illawarra Area Health Service–Eastern Section Human Research Ethics Committee. After participants gave informed consent, the following measures were administered to participants in an individual testing session.

Table 1. Demographic characteristics of the control, mild cognitive impairment (MCI), and dementia participants.

	Control (<i>n</i> = 53)		MCI (<i>n</i> = 48)		Dementia (<i>n</i> = 39)	
	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>	<i>M</i>	<i>SD</i>
<i>Participant characteristics</i>						
Age (years)	77.8	4.66	78.6	4.87	79.8	6.19
Education (years)	11.3	3.28	12.2	3.90	12.03	4.46
Gender (% male)	41.5		54.2		51.3	
MMSE	28.7	1.42	28.0	1.56	25.3	4.30
<i>Prospective memory</i>						
VW number correct (maximum = 8)	3.58	2.05	2.00	2.04	0.97	1.29
<i>Retrospective memory</i>						
VW tasks recalled	5.40	1.53	4.32	1.76	3.09	2.18
VW tasks recognized	7.54	0.81	7.04	0.93	6.00	2.05
<i>Cognitive functioning</i>						
Visual Span	11.86	3.58	9.96	3.57	8.10	3.92
TOL (Excess moves)	7.39	5.47	8.33	6.11	16.91	18.04

Note. MMSE refers to the Mini-Mental State Examination; VW refers to Virtual Week; TOL refers to Tower of London.

Virtual Week

Performance on an adapted version of this laboratory measure of PM was the primary dependent measure of interest. As noted, Virtual Week was selected as it has been consistently identified as a reliable indicator of PM (for a review, see Rendell & Henry, 2009), and the incorporation of the different types of PM tasks means that PM performance can be investigated systematically in relation to different PM task demands.

Virtual Week is a computerized board game, in which participants move around the board with the roll of a dice. The times of day people are typically awake are marked on the board, with each circuit of the board representing a day. As participants move around the board a series of events occur, in which a screen pops up describing an event (e.g., “you stop at a café for lunch”), the participant is required to make choices about the event (e.g., what to eat) and in some of these events remember to carry out an event-based PM task (e.g., take medication). The original version (Rendell & Craik, 2000) was modified in the present study for older adults with MCI and early stages of dementia, by reducing the overall task requirements to 4 (instead of 10) PM tasks per virtual day, over 2 (instead of 7) simulated days (plus a practice day). Two PM tasks were time-based (i.e., triggered by passing a particular time on the board), and two were event-based (i.e., triggered by encountering a specific event in the game). For each virtual day, one of the time-based and one of the event-based tasks was regular (a routine task involving remembering to take medication which recurred every day), and one of the time-based and one of the event-based tasks was irregular (a one-off, nonrecurring task that differed for each virtual day). Virtual Week was administered on an HP Tablet Notebook TC4400 via a touch-screen computer interface. The Virtual Week gameboard, cards, and dice were

electronically presented, and the participants interacted with the game by tapping the screen with a pen-stylus. As in Rendell and Craik (2000), participants were given pre-game instructions and then a practice virtual day to ensure they understood all features of the game, thus regular tasks occurred three times and irregular tasks only once.

Responses were scored as the number (out of eight) *Correct*, *Missed*, *Wrong*, *Late*, *Early*, *Little Late*, *Little Early*, or *Cancel*. *Correct* scores indicated the target item was remembered at the correct time (correct time was after the dice roll for the move that took the token onto or past the target square and before the next roll of the dice); participants were marked *Wrong* when they selected the wrong task; *Missed* indicated the participant did not remember the target item at any time; *Little Late* items were remembered after the correct time criterion but within two further rolls of the dice and *Late* items were after the *Little Late* criterion and before the end of the virtual day. *Little Early* and *Early* items were the converse of *Late* items; *Little Early* was within two dice rolls before the correct time criterion and *Early* was before the *Little Early* criterion and after the start of the virtual day. *Cancel* was when participants opened the perform task list and closed the list without selecting a task. In the present study, Cronbach’s alpha for the total score was estimated to be .74.

Retrospective memory

After completion of the virtual days, the participant was asked to recall the PM tasks. Once the participant could recall no more tasks, a list of randomly ordered correct tasks and foils was given to the participant to check off the tasks that were in the game. This provided a measure of both recall and recognition of the tasks. Cronbach’s alpha was estimated to be .65 for recall and .69 for recognition.

Working memory

The Colorado Assessment Tests (CATS) Visual Span task (Davis & Keller, 2002) is a computerized version of a block tapping task (Corsi, 1972) used to assess working memory. It is analogous to established digit-span testing (Wechsler, 1997), with spans of increasing length being presented, and recalled by the examinee in a forward and then backward direction. Normative data shows increasing performance with age until the twenties and then declining performance thereafter (Davis & Keller, 2002).

Executive functioning

The Tower of London test (Davis & Keller, 2002; Shallice, 1982) is used to assess executive control, and specifically planning and execution skills, which are considered to be particularly relevant to PM function (Phillips, Henry, & Martin, 2008). The CATS Tower of London (Davis & Keller, 2002) is a computerized version of the task which involves moving different colored pegs to match an arrangement shown in a target picture. The CATS version includes trials of varying complexity, thereby overcoming ceiling effects seen in previous versions of the Tower of London (Tunstall, 1999). Normative data for the CATS version show increasing performance with age until the twenties and then declining performance thereafter (Davis & Keller, 2002). Studies using a similar (noncomputerized) version of the task have reported good reliability and criterion-related validity (Culbertson & Zillmer, 1998).

RESULTS

Background Cognitive Measures

There were group differences in Visual Span, $F(2,134) = 17.01$; $p < .001$; $\eta_p^2 = .20$; and the Tower of London, $F(2,131) = 9.34$; $p < .001$; $\eta_p^2 = .13$ (see Table 1). Follow-up Tukey HSD tests indicated that both measures differentiated between dementia and controls (both $ps < .001$). Visual Span differentiated between the control and MCI groups ($p = .016$), and between the MCI and dementia groups ($p = .006$). Tower of London differentiated between the MCI and dementia groups ($p = .002$), but not the MCI and control groups ($p = .770$).

Prospective Memory: Virtual Week

The number of correct PM responses are presented in Figure 1 as a function of *group* (control, MCI, dementia) and *PM cue* (time, event). These data were analyzed with a $3 \times 2 \times 2$ mixed analysis of variance with the between-subjects variable of *group* and the within-subjects variables of *PM task* (regular, irregular) and *PM cue* (time, event). There was no main effect of PM task, $F(1,137) = 0.42$; $p = .520$; $\eta_p^2 < .01$, but there was a main effect of PM cue $F(1,137) = 24.90$; $p < .001$; $\eta_p^2 < .15$ and of group, $F(2,137) = 23.32$, $p < .001$, $\eta_p^2 = .25$. None of the two- or three-way interactions were significant (all F s < 1.2 , $ps > .320$). The main effect of PM cue indicated better performance in response to time-based than

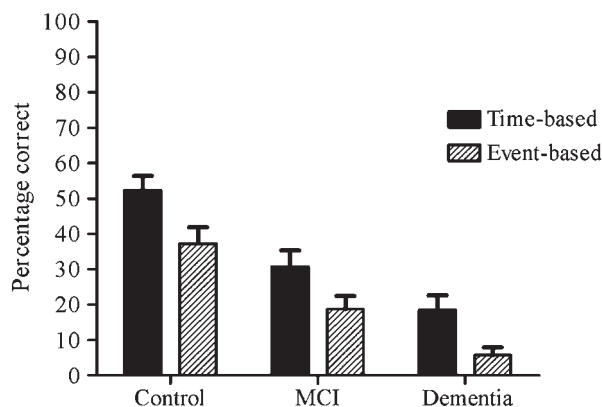


Fig. 1. Number of correct responses on the Virtual Week as a function of prospective memory cue type (time & event) for controls and participants with mild cognitive impairment (MCI) and dementia. Bars represent SE.

event-based cues. Follow-up Tukey HSD tests of the group main effect indicated that the dementia and MCI groups were each impaired relative to the control group (both $p < .001$) and the dementia group in turn performed more poorly than the MCI group ($p = .035$). There was no difference in PM performance between MCI subtypes, $F(2,45) = .583$; $p = .56$; $\eta_p^2 = .03$. Nor was there any difference between the amnesic MCI subtypes and non-amnesic MCI subtype, $t(46) = .284$; $p = .78$; $\eta_p^2 = .03$.

Figure 2 shows the pattern of errors on Virtual Week. Most of the errors involved a failure to respond (missed responses), with all other error types relatively infrequent. Thus the PM tasks were nearly always either remembered reasonably accurately or not remembered at all.

Differentiating the Prospective and Retrospective Components of Virtual Week

The next step in the analyses involved assessing whether any of the group differences observed on Virtual Week could be attributed to difficulties with the retrospective memory component of the task (i.e., correctly recalling the tasks that needed to be done). Performance on this component was measured through recall and recognition tests of the eight Virtual Week tasks (see Table 1). The results indicated a main effect of group for total number of items correctly recalled, $F(2,128) = 16.86$; $p < .001$; $\eta_p^2 = .21$; and recognized, $F(2,128) = 11.20$; $p < .001$; $\eta_p^2 = .15$. Although there was no main effect of PM task, a follow-up t test comparing percentage *recall* of regular and irregular tasks indicated that of the two task types, regular tasks were more likely to be recalled following completion of Virtual Week, $t(129) = 10.66$; $p < .001$; $\eta_p^2 = .47$. These data are consistent with earlier studies showing that irregular (relative to regular) tasks impose greater demands on retrospective memory. To explore the patterns of shared variance in Virtual Week, two analyses of covariance (ANCOVA) were conducted with the dependent variable of PM Total Score and either total number of Virtual Week items (i) recalled or (ii) recognized entered as



Fig. 2. Types of error responses on Virtual Week for controls and participants with mild cognitive impairment (MCI) and dementia.

covariate. Following entry of each covariate, the group effect size (η_p^2) was reduced from .25 to .10 and .18, respectively, but remained significant ($ps = .002$ and $< .001$, respectively). Consequently, although interpretation of ANCOVA designs requires caution (Miller & Chapman, 2001), these data suggest that difficulties with the retrospective component of the task contributed to the group effects observed on Virtual Week, but significant residual variance is attributable to a separable prospective component. Consistent with this interpretation, the correlation between Virtual Week and group status remained significant after partialling out the number of Virtual Week items correctly recalled ($r_p = .28$; $p = .001$) and recognized ($r_p = .41$; $p < .001$).

Cognitive Correlates of Prospective Memory Function

Group effects were observed on the measures of Visual Span and Tower of London. Correlational analyses indicated that Virtual Week performance was related to both these measures ($rs = .49$ and $.27$, respectively; both $ps < .01$). To explore the patterns of shared variance in the dataset, further exploratory ANCOVAs were conducted, this time with the dependent variable of PM Total Score, and one of the cognitive measures entered as a covariate. After exploring the influence of Visual Span and Tower of London as covariates individually, the final ANCOVA entered both measures as covariates simultaneously. The group effect size (η_p^2) was reduced from .25 to .13, .20 and .12, respectively, but for all ANCOVAs remained significant (all $ps < .05$). Again, consistent with this interpretation, the correlation between Virtual Week and group status remained significant after partialling out Visual Span ($r_p = .35$; $p < .001$), Tower of London ($r_p = .44$; $p < .001$), and both cognitive measures simultaneously ($r_p = .34$; $p < .001$).

DISCUSSION

These data confirm previous research studies in showing that individuals with MCI and dementia exhibit PM difficulties

relative to demographically matched controls. However, because prior studies have failed to agree on whether reliable group differences exist on measures of this construct between MCI and dementia (Kazui et al., 2005; Troyer & Murphy, 2007), the present results are important in showing that individuals with dementia are more impaired on Virtual Week than those with MCI. These data affirm the importance of using measures of PM that have documented reliability and validity. As noted previously, Virtual Week has been well validated, and shown to be sensitive to age-related cognitive impairment specifically (Rendell & Henry, 2009; Will et al., 2008). It seems likely that the discrepancies in prior studies reflected method variance, and in particular, fewer opportunities given to perform the PM task in the study by Kazui et al. (2005) (three) relative to the study by Troyer and Murphy (2007) (eight). As in the latter study, the present study afforded participants eight opportunities to execute the PM task.

The second aim of the study was to assess whether individuals with MCI and dementia exhibit a differentiated profile of impairment on Virtual Week. In line with prior research results, it was predicted that PM tasks which imposed relatively greater demands on strategic resources may be particularly sensitive to the presence of MCI, although they may not differ in their relative sensitivity to dementia. In fact no interaction between group and either cue type (time, event) or task type (regular, irregular) was observed, indicating that both clinical groups exhibited a relatively generalized level of impairment on Virtual Week that did not vary as a function of specific task demands.

In the PM literature, most weight has typically been attributed to the distinction between time- and event-based cues. Prior studies that have manipulated cue-type in this group have shown individuals with MCI to be disproportionately impaired when responding to time-based cues (Karantzoulis et al., 2009; Troyer & Murphy, 2007) but those with dementia to be equivalently impaired (Troyer & Murphy, 2007). It is suggested that the absence of an interaction with cue type in the present study may reflect the unique manner in which Virtual Week operationalizes these parameters. Relative to most laboratory time-based PM tasks, the time-based tasks

in Virtual Week have considerable external cues (the time is cued by the activities relevant to the virtual time of day and the time is clearly seen and “encountered” on the Virtual Week Board Game). The provision of these cues may, therefore, have equated these tasks to the event-based tasks in terms of their reliance on self-initiated processing, representing the situation in daily life where some times of day can have strong environmental cues. Consequently, we do not wish to over-interpret the absence of interaction with cue-type identified in the present study.

More compelling was the absence of any interaction between group and task regularity on Virtual Week. Previous research has shown that the regular tasks in Virtual Week require fewer processing resources (and indeed, in the present study a *post hoc* analysis of task recall following completion of Virtual Week also suggested that the regular tasks imposed fewer demands on retrospective memory—i.e., irregular, relative to regular, tasks were less likely to be successfully recalled retrospectively). The absence of any interaction effect, therefore, suggests that the MCI- and dementia-related difficulties observed on Virtual Week may not simply be restricted to the retrospective memory component, but also extend to the PM component (i.e., the implementation of delayed intentions). This interpretation is also consistent with the analyses in which number of Virtual Week items correctly recalled and recognized were covaried. While the use of ANCOVA in nonrandomized designs has been subject to some debate, it has been suggested that this methodology may be useful (despite nonrandom assignment) in the context of exploration of a dataset to understand patterns of shared variance (Huitema, 1980; Miller & Chapman, 2001). Although speculative, taken together these analyses suggest that while difficulties with the retrospective component of the task (as well as working memory and executive functioning) may each have contributed to the group effects observed on Virtual Week, significant residual variance may be attributable to a separable, prospective component. However, given the noted difficulties of ANCOVA use in nonrandomized designs, coupled with the use of single indicators to tap the key cognitive constructs of interest, clearly further research is needed to test this interpretation of these data.

A strength of this study is the recruitment of participants diagnosed by use of consistent diagnostic criteria for MCI and dementia; however, some limitations must be acknowledged. Some of the dementia group were taking dementia medication, but it was not possible to conduct formal analyses assessing medication status and cognitive performance due to the very differing medication types and dosages participants received. The role of medication status in PM function in this group, therefore, remains an important issue for future research. Given that the purpose of such medication is to improve cognition, one possibility is that use of such medication may serve to attenuate dementia effects on PM.

As noted previously, the present study used a modified version of Virtual Week, in which task demands were reduced. Despite these modifications some participants, particularly those with dementia, performed at zero. Thus, while

a key strength of Virtual Week is its sensitivity to even very early signs of cognitive decline, further modifications which decreased task difficulty would be needed to further extend its usefulness as a research tool in this particular population.

In conclusion, it has long been recognized that deficits in the ability to implement delayed intentions are likely to lead to problems in daily functioning. The current study indicates that both MCI and dementia are associated with PM deficits, and supports the use of Virtual Week in clinical practice as a tool to quantify the magnitude of these impairments. It is suggested that while failures of retrospective memory, working memory, and executive functioning each potentially contribute to PM difficulties in each of these groups, they may not be sufficient to account for the magnitude of the PM impairment observed.

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REFERENCES

- Albert, M.S., & Blacker, D. (2006). Mild cognitive impairment and dementia. *Annual Review of Clinical Psychology*, 2, 379–388.
- Artero, S., Petersen, R., Touchon, J., & Ritchie, K. (2006). Revised criteria for mild cognitive impairment: Validation within a longitudinal population study. *Dementia and Geriatric Cognitive Disorders*, 22, 465–470.
- Barnes, J., Ourselin, S., & Fox, N.C. (2009). Clinical application of measurement of hippocampal atrophy in degenerative dementias. *Hippocampus*, 19, 510–516.
- Blanco-Campal, A., Coen, R.F., Lawlor, B.A., Walsh, J.B., & Burke, T.E. (2009). Detection of prospective memory deficits in mild cognitive impairment of suspected Alzheimer’s disease etiology using a novel event-based prospective memory task. *Journal of the International Neuropsychological Society*, 15, 154–159.
- Chasteen, A.L., Park, D.C., & Schwarz, N. (2001). Implementation intentions and facilitation of prospective memory. *Psychological Science*, 12, 457–461.
- Corsi, P.M. (1972). Human memory and the medial temporal region of the brain. *Dissertation. Abstracts International*, 34.
- Culbertson, W.C., & Zillmer, E.A. (1998). The construct validity of the Tower of London as a measure of the executive functioning of ADHD children. *Assessment*, 5, 215–226.
- Davis, H.P., & Keller, F. (2002). *Colorado Assessment Tests (CATS), version 1.2*. Colorado Springs, CO.
- den Ouden, H.E., Frith, U., Frith, C., & Blakemore, S.J. (2005). Thinking about intentions. *Neuroimage*, 28, 787–796.
- Duchek, J.M., Balota, D.A., & Cortese, M. (2006). Prospective memory and Apolipoprotein E in healthy aging and early stage Alzheimer’s disease. *Neuropsychology*, 20, 633–644.

- Glisky, E.L. (1996). Prospective memory and the frontal lobes. In M.A. Brandimonte, G.O. Einstein, & M.A. McDaniel (Eds.), *Prospective Memory: Theory and Applications* (pp. 249–266). Mahwah, NJ: Lawrence Erlbaum Associates.
- Henry, J.D., Macleod, M.S., Phillips, L.H., & Crawford, J.R. (2004). A meta-analytic review of prospective memory and aging. *Psychology and Aging, 19*, 27–39.
- Huitema, B. (1980). *Analysis of Covariance and Alternatives*. New York: Wiley.
- Jones, S., Livner, A., & Backman, L. (2006). Patterns of prospective and retrospective memory impairment in preclinical Alzheimer's disease. *Neuropsychology, 20*, 144–152.
- Karantzoulis, S., Troyer, A.K., & Rich, J.B. (2009). Prospective memory in amnesic mild cognitive impairment. *Journal of the International Neuropsychological Society, 15*, 407–415.
- Kazui, H., Matsuda, A., Hirono, N., Mori, E., Miyoshi, N., Ogino, A., et al. (2005). Everyday memory impairment of patients with mild cognitive impairment. *Dementia and Geriatric Cognitive Disorders, 19*, 331–337.
- Kinsella, G.J., Ong, B., Storey, E., Wallace, J., & Hester, R. (2007). Elaborated spaced-retrieval and prospective memory in mild Alzheimer's disease. *Neuropsychological Rehabilitation, 17*, 688–706.
- Kliegel, M., Martin, M., McDaniel, M.A., & Einstein, G.O. (2002). Prospective memory and aging: How planning affects performance. Paper presented at the Cognitive Aging Conference, Atlanta, GA, USA.
- Kochan, N., Brodaty, H., Crawford, J., Slavin, M., Low, L., Trollor, J., et al. (2009). *How to define the cognitive impairment in Mild Cognitive Impairment: Comparison of different neuropsychological classification methods using data from the Sydney Memory and Ageing Study*. Paper presented at the International Conference on Alzheimer's Disease, Vienna, Austria.
- Martin, M., Kliegel, M., & McDaniel, M.A. (2003). The involvement of executive functions in prospective memory performance of adults. *International Journal of Psychology, 38*, 195–206.
- Martins, S.P., & Damasceno, B.P. (2008). Prospective and retrospective memory in mild Alzheimer's disease. *Arquivos de Neuropsiquiatria, 66*, 318–322.
- McDaniel, M.A., & Einstein, G.O. (1992). Aging and prospective memory: Basic findings and practical applications. *Advances in Learning and Behavioral Disabilities, 7*, 87–105.
- McDaniel, M.A., & Einstein, G.O. (2007). *Prospective Memory: An Overview and Synthesis of an Emerging Field*. Thousand Oaks, CA: Sage Publications, Inc.
- Miller, G.A., & Chapman, J.P. (2001). Misunderstanding analysis of covariance. *Journal of Abnormal Psychology, 110*, 40–48.
- Petersen, R.C. (2007). Mild cognitive impairment: Current research and clinical implications. *Seminars in Neurology, 27*, 22–31.
- Phillips, L.H., Henry, J.D., & Martin, M. (2008). Adult aging and prospective memory: The importance of ecological validity. In M. Kliegel, M.A. McDaniel, & G.O. Einstein (Eds.), *Prospective Memory: Cognitive, Neuroscience, Developmental, and Applied Perspectives*. Mahwah: Erlbaum.
- Rendell, P.G., & Craik, F.I.M. (2000). Virtual week and actual week: Age-related differences in prospective memory. *Applied Cognitive Psychology, 14*, S43–S62.
- Rendell, P.G., & Henry, J.D. (2009). A review of Virtual Week for prospective memory assessment: Clinical implications. *Brain Impairment, 10*, 14–22.
- Reynolds, J.R., West, R., & Braver, T. (2009). Distinct neural circuits support transient and sustained processes in prospective memory and working memory. *Cerebral Cortex, 19*, 1208–1221.
- Scheltens, P. (2009). Imaging in Alzheimer's disease. *Dialogues in Clinical Neuroscience, 11*, 191–199.
- Schmitter-Edgecombe, M., Woo, E., & Greeley, D.R. (2009). Characterizing multiple memory deficits and their relation to everyday functioning in individuals with mild cognitive impairment. *Neuropsychology, 23*, 168–177.
- Shallice, T. (1982). Specific impairments of planning. *Philosophical Transactions of the Royal Society of London. Series B, Biological Sciences, 298*, 199–209.
- Smith, G., Della Sala, S., Logie, R.H., & Maylor, E.A. (2000). Prospective and retrospective memory in normal ageing and dementia: A questionnaire study. *Memory, 8*, 311–321.
- Troyer, A.K., & Murphy, K.J. (2007). Memory for intentions in amnesic mild cognitive impairment: Time- and event-based prospective memory. *Journal of the International Neuropsychological Society, 13*, 365–369.
- Tunstall, J. (1999). *Improving the Utility of the Tower of London: A Neuropsychological Test of Planning*. Brisbane, Australia: Griffith University.
- Wechsler, D. (1997). *Wechsler Adult Intelligence Scale - III*. San Antonio: The Psychological Corporation.
- Will, C.M., Rendell, P.G., Ozgis, S., Pierson, J.M., Ong, B., & Henry, J.D. (2008). Cognitively impaired older adults exhibit comparable difficulties on naturalistic and laboratory prospective memory tasks. *Applied Cognitive Psychology*. Retrieved from (www.interscience.wiley.com) DOI: 10.1002/acp.1514.
- Wilson, B., Cockburn, J., Baddeley, A., & Hiorns, R. (1989). The development and validation of a test battery for detecting and monitoring everyday memory problems. *Journal of Clinical and Experimental Neuropsychology, 11*, 855–870.
- Zöllig, J., West, R., Martin, M., Altgassen, M., Lemke, U., & Kliegel, M. (2007). Neural correlates of prospective memory across the lifespan. *Neuropsychologia, 45*, 3299–3314.