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A COMPARISON OF LABORATORY, CLINICAL AND SELF-REPORT MEASURES OF  
PROSPECTIVE MEMORY IN HEALTHY ADULTS AND INDIVIDUALS WITH BRAIN  
INJURY

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## Abstract

Individuals with traumatic brain injury (TBI) have demonstrated deficits in prospective memory (PM) functioning when compared to healthy adults. These deficits have been measured using laboratory measures, clinical measures and self-report questionnaires. However, PM has been shown to involve multiple cognitive processes and a variety of stages. Thus, it is not known if these measures all assess the same aspects of PM. This study used all three types of measures in healthy adults and individuals with TBI to determine which aspects of PM were affected. Results demonstrated the sensitivity of all three types of measures to PM deficits in TBI. Data from the laboratory measure suggested a specific difficulty with detecting the correct cue. Data from the clinical measure suggested that TBI has a greater effect on time-based cues than event-based cues and that the primary deficit is a prospective intention retrieval deficit rather than the retrospective memory component. In addition, those with TBI did not differ from healthy adults when the time delay was short enough, suggesting that PM is not universally impaired. Data from the self-report questionnaire suggested that those with TBI are more sensitive to difficulties with basic activities of daily living rather than instrumental activities on daily living. These results are discussed in terms of rehabilitation techniques that could focus first on cue detection and use basic activities of daily living as outcome measures.

## Introduction

Prospective memory (PM), remembering to remember and carry out a future action, is an important part of everyday life (Ellis & Freeman, 2008; Rendell & Thompson, 1999). For example, remembering to take out the trash or take medications at the correct time are both everyday PM tasks. For individuals with neurological disorders, including traumatic brain injury (TBI), schizophrenia and Parkinson's disease, PM deficits have been shown to impact daily functioning (Mathias & Mansfield, 2005; Raskin, 2009; Raskin et al., 2012; Raskin et al., 2014; Shum, Valentine & Cutmore, 1999; Shum, Levin & Chan, 2011; Shum et al., 2004).

PM has been conceptualized using two current theories. The multi-process view of PM suggests that PM performance can be either spontaneous or rely on strategic monitoring, depending on the nature of the task. Differences can occur due to disparities in type of task, type of cue, the nature of the ongoing task, and the individual. Thus, for example, tasks that must be remembered at a particular time have been suggested to require more strategic monitoring than those that must occur in response to an external cue (e.g., Einstein, McDaniel, Richardson, Guynn, & Cunfer, 1995). More recently Scullin, McDaniel, & Shelton (2013) expanded this model to suggest there is a dynamic interplay between the spontaneous retrieval and the strategic monitoring, such that individuals will monitor in a context where cues are expected but then disengage from monitoring when cues are not expected. They refer to this as the Dynamic Multiprocess Framework. The preparatory attentional and memory processes (PAM) theory, on the other hand, asserts that PM always requires attentional processes that are resource demanding due to explicit monitoring or the need to maintain the intention (Smith, 2003). This is assumed to be the PM aspect of the task whereas the recall of what one was supposed to do is based on retrospective memory (RM). Given the prevalence of attention deficits in populations such as

those with TBI, and the importance of PM performance to everyday life, investigating whether some aspects of PM might be spared could lead to important directions in management post TBI.

Measures of PM all have some common elements. These include the encoding and formation of an intention to be performed in the future, an ongoing task during the delay period, and a cue to signal it is time to perform the intention. Research on this topic has been performed using laboratory, self-report and clinical measures. It is thought that laboratory measures provide greater ability to manipulate specific variables to answer particular theoretical questions (e.g., Einstein & McDaniel, 2005). Self-report measures provide data on the subjective experience of PM (Roche et al., 2007) whereas clinical measures have been used in attempts to approximate more real-world applications of PM, such as calling the experimenter at a specified time (Rendell & Thomson, 1999). Each of these measures provides valuable data on PM performance, but clear and consistent relationships between them have yet to be defined (e.g., Raskin & Sohlberg, 2009; Rendell & Thompson, 1999; Uttl & Kibreab, 2011). Given the possibility that these measures are all capturing unique and not entirely overlapping aspects of PM, it is important to compare the findings from each.

Einstein and McDaniel (1990) introduced a dual-task laboratory paradigm that has been widely used in the literature. Like all PM tasks, there is a prospective remembering task embedded in an ongoing task, followed by a delay period. After this period, the ongoing task is reintroduced without a reminder of the PM task. The PM task, embedded in the ongoing task, is presented and the response of the participant is measured (Einstein & McDaniel, 2005). This laboratory paradigm has proven invaluable in testing specific theoretical questions about PM performance. However, these paradigms tend to involve only one type of PM response performed repeatedly, and the ecological validity is considered low (Delprado, Kinsella, Ong,

Pike, Ames, Storey, Saling, Clare, Mullaly, & Rand, 2012). In addition, the sensitivity of these paradigms to neurological disorders appears to vary with the characteristics of the PM and ongoing tasks (e.g., Foster, McDaniel, Repovs, & Hershey, 2009).

There are also a number of currently used self-report PM measures, including the Comprehensive Assessment of Prospective Memory (CAPM; Chau, Lee, Fleming, Roche, & Shum, 2007), the Prospective and Retrospective Memory Questionnaire (PRMQ, Crawford, Smith, Maylor, Della Sala, & Logie, 2003), and the Prospective Memory Questionnaire (Hannon, Adams, Harrington, Fries-Dias, & Gipson, 1995). Essentially, these measures all quantify common PM tasks that are undertaken in daily life and as such may have a higher personal significance than the other measures. Questionnaires have arguably high ecological validity and allow for a greater understanding of the experience of PM failures in daily life. They are generally brief, easy to use and understand, and time-efficient (Man et al., 2011). However, they have not been well-correlated with other measures of PM, perhaps due to limited insight or awareness in individuals with brain injury (Fleming et al., 2009; Roche et al., 2002).

The CAPM is one questionnaire that was designed to measure PM failures. It has been demonstrated to have good reliability (Chau, Lee, Fleming, Roche, & Shum, 2007). The full CAPM contains three sections, one that measures frequency of PM failure, one that measures the perceived importance of these failures and the third that measures the perceived reasons for forgetting. It can be completed by an individual or by a family member. A principal components analysis of the first part, Section A, revealed that there were two components, one that related to basic activities of daily living (BADL), such as daily self-care, and one that related to instrumental activities of daily living (IADL), such as household management activities

(Chau, et al., 2007). The CAPM has been used in previous studies with individuals who have TBI and has demonstrated sensitivity to perceived failures in this population (Huang et al., 2014) and has been analyzed according to Ellis's (1996) five phases of prospective memory to demonstrate that those with TBI had difficulty with encoding and formation and initiation of prospective memories (Roche, Moody, Szabo, Fleming & Shum, 2007).

To our knowledge, there are currently four standardized clinical measures of PM and these are considered more naturalistic than laboratory measures. These are the Memory for Intentions Test (MIST) (Raskin, Buckheit, & Sherrod, 2010), the Cambridge Assessment of Prospective Memory Test (CAMPROMPT) (Wilson et al., 2005), Virtual Week (Rendell & Henry, 2009) and the Royal Prince Alfred Memory Test (Radford, Lah, Say, & Miller, 2011). These tests have all shown good psychometric properties and sensitivity to neurological disorders.

Of these, the MIST has the ability to separate out different aspects of performance, such as type of cue and time delay. The MIST includes both time-based and event-based cues that are either two minutes or fifteen minutes in duration between encoding and performance. The time-based cues allow for self-initiated retrieval ("In fifteen minutes, tell me that it is time to take a break") and, unlike most laboratory tasks, the event-based cues are related to the events that need to be performed (i.e. "When I hand you a red pen, sign your name on the paper"). The response of participants can either be an action or verbal response. Due to the MIST's ability to measure different attributes of PM, it also allows for six types of errors to be analyzed if failure of a PM task were to occur (Raskin, 2009). The MIST has been demonstrated to be sensitive to PM deficits in individuals with TBI (Raskin, 2009), Parkinson's disease (Raskin et al., 2011), HIV

(e.g., Woods et al., 2010), and schizophrenia (Raskin et al., 2014; Twamley et al., 2007) and has been shown to have good psychometric properties (Raskin, 2009; Woods, et al., 2008).

Previous studies comparing questionnaires and clinical measures have yielded mixed results. One previous study found no correlation between the CAPM and total score on the MIST, suggesting that either the two measures are measuring separate functions or that subjective experience and objective performance are not the same (Fleming et al., 2009). Relatives' report on frequency of PM failure did correlate with the MIST total score as well as both the time-based cue and event-based cue scales, which suggest that the difference may be due to poor subjective awareness. No data were provided on the relationship between the CAPM and the 24 hour delay item on the MIST. Another study used the PRMQ and found that while the MIST did not correlate with responses on the self-report questionnaire, the MIST did correlate with performance in daily life on 10 items over a week as reported by a significant other (Raskin & Sohlberg, 2009). In a study of healthy older adults the MIST was uniquely related to the IADL component of the PRMQ (Woods et al., 2012). Thus, this again suggests that the questionnaires are tapping into an aspect of the individual's perception of his/her performance that may not be related to the individual's actual performance.

It seems possible, and even likely, that these three types of measures are sampling different attributes of PM. For example, it has been argued that the cognitive demands in daily life, as measured by a questionnaire, may be very different than those in a laboratory setting (Fleming, Doig, & Katz, 2000). As has been suggested, different PM tasks could very well challenge different cognitive systems depending on the task (McDaniel & Einstein, 2011). In fact, there is growing evidence that different types of PM tasks rely on different brain systems. For example, tasks with focal cues may require little strategic monitoring and thus rely on the



medial temporal structures while those with non-focal cues may require activity of the prefrontal cortex (e.g., McDaniel & Einstein, 2011).

These tasks may also require different levels of attentional resources. The laboratory measures tend to use only event-based cues, thus, they may be less attention demanding than the time-based tasks in the MIST. Furthermore, unlike the MIST, the event-based cue in the laboratory measure is usually embedded within the ongoing task, which may further serve to focus attention on the cue. On the other hand, in the laboratory paradigm the embedded PM task (i.e. press a key when you see a specific word) does not explicitly put the participant in a “retrieval mode” and instead requires participants to enter this mode on their own. In contrast, the event cues of the MIST puts participants in a retrieval mode, (i.e. by handing the participants a form requesting medical records) in which they may be aware something should happen because of the high association between the intended action and the cue. On the other hand, the CAPM queries performance on both event-based (e.g., forgetting to lock the door when leaving home) and time-based (e.g., forgetting to take tablets at a prescribed time) everyday tasks. It can be argued that the CAPM is the only one that measures tasks in which the individual must form his/her own intention and create his/her own plan for execution (e.g., Dobbs & Reeves, 1996). For the other tasks this is provided for the participants by the experimenter. However, because the CAPM relies on retrospective subjective report, it is not known if it is an accurate reflection of performance.

Differences between the measures may shed light on research findings based on clinical populations. Research on TBI has revealed significant deficits in PM performance, presumably due to impairment of the prefrontal cortical regions, and individuals with specific frontal lobe damage have shown impairments in PM (Neulinger, Oram, Tinson, O’Gorman & Shum, 2015).

These individuals show deficits in the total score on the MIST as well as differences in performance on the two- and fifteen-minute time delays (Raskin, 2009). Adults with TBI, like healthy adults, are better at event-based cues, than at time-based cues (Mioni, Rendell, Henry, Cantagallo, & Stablum, 2013; Raskin, 2009). Studies on participants with TBI using the laboratory paradigms have also shown difference in performance. In a study done by Shum, Valentine, & Cutmore (1999) using both time-, event-, and activity-based PM tasks. The results revealed differences in all three tasks, with the individuals with TBI performing significantly worse on all tasks (Shum et al. 1999).

In reviewing the literature, however, we are not aware of a prior study that used all three types of measures with individuals with TBI. Thus, it would be worthwhile to measure all three of them within the same population. In the first place, this would allow for a more comprehensive evaluation of both laboratory-based and naturalistic PM functioning. Second, it would allow for an analysis of the specific aspects of PM being measured by each task.

This study compared performance of healthy adults and individuals with TBI on the MIST (Raskin, 2009), an Einstein and McDaniel dual-task laboratory paradigm that was previously published by Pereira, Ellis and Freeman (2012), and the CAPM (Chau et al., 2007). It aimed to understand how individuals with TBI perform on each measure given concurrently. The Pereira et al. (2012) task also allows for an analysis of the effect of cue-action relatedness. The results of this study would provide insight as to whether the three measures are measuring the same or different aspects of PM. In addition, a Stroop Color-Word Interference task was administered within the laboratory measure to evaluate if there is any effect of poor executive functioning on PM performance.

## **Methods**

*Participants:*

One hundred healthy adults (51 females) and 50 people with TBI (18 females) took part in this study. Demographic information for the two groups is presented in Table 1. There were no significant differences between the groups on any demographic variables.

*Inclusion criteria:* All participants spoke English as a primary language. Individuals with TBI were at least one-year post injury, had lowest postresuscitation Glasgow Coma Score (Teasdale & Jennett, 1974) of 8-12, and had obtainable medical records. People with TBI were recruited through local hospitals and the local brain injury association in the US. Healthy adults were relatives of the people with brain injury, employees of Trinity College, or members of the community. Descriptive injury-related information of the TBI group is presented in Table 2.

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Insert Tables 1 and 2 about here

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*Exclusion criteria:* Any potential participant that had previous neurological or psychiatric illness, history of substance abuse or diagnosed learning disability, visual impairment that would interfere with reading the test materials, and/or experienced seizure during the previous year was excluded from the study.

*Screening measures:* All participants were administered the Brief Psychiatric Rating Scale (Overall and Gorham, 1962) to screen for psychiatric illness. No individual with severe depression ( $\geq 21$  on the Beck Depression Inventory) (Beck, 1987) anxiety (Beck Anxiety

Inventory) ( $\geq 30$  on the Beck, 1990), or global cognitive dysfunction (severe impairment on four or more of the subscales of the Neurobehavioral Cognitive Status Examination) (Kiernan, Mueller, Langston, & Van Dyke, 1987) was included.

*Measures:*

The Memory for Intentions Test (MIST)

The MIST (Raskin, 2010) is a 30-min, 8-trial test during which participants engage in a word search puzzle as the ongoing task. A complete description of the MIST administration and scoring procedures can be found in Raskin (2009). Briefly, it is comprised of four trials with event-based cues (e.g., “When I hand you a postcard, self-address it.”) and four trials with time-based cues (e.g., “In 15 minutes, tell me it is time to take a break.”), with each item scored from 0-2 points; thus, the separate event-based and time-based scales have scores ranging from 0 to 8. The time- and event-based trials are balanced for delay interval (i.e., 2- and 15-min delay periods) and response modality (i.e., verbal and action responses). The MIST allows for separate scoring of time-based trials (8 points possible), event-based trials (8 points possible), 2-min delay periods (8 points possible), 15-min delay periods (8 points possible), verbal response trials (8 points possible) and action response trials (8 points possible), which are summed for a total of 48 possible points. However, this involves inclusion of the score of each trial three times in the total score (e.g., Trial 1 is a 2-minute delay trial, time-based cue, and verbal response, thus contributing to the 2-minute delay, time-based cue, and verbal response scores). A large digital clock is in full view of the participant at all times. For the event-based trials, the cues all have high cue-action relatedness and were considered to be ecologically relevant, meaning they are related to the response required and could naturally elicit that required response (e.g., When I hand you a request for records form, please write your doctors’ names on it). The ongoing task

is non-focal as the word search is not related to the PM items. Prior studies support the reliability (Raskin, 2009; Woods et al., 2008) and construct validity (e.g., Raskin, Buckheit, & Sherrod, 2010; Woods et al., 2009) of the MIST.

At the completion of the eight MIST trials, participants are given eight multiple choice recognition items (e.g., “At any time during this test, were you supposed to: 1) tell me to make an appointment; 2) tell me when I can call you tomorrow; 3) tell me to call for a prescription.”). The recognition scale is included as a way to determine whether PM failures are due to encoding versus retrieval failures. Impairment on recognition items is likely to reflect deficits in RM rather than PM functions. Furthermore, a 24-hr delay trial was administered for which examinees were instructed to leave a voicemail message for the examiner the day after the exam indicating the number of hours the participant slept the night after the evaluation. In addition, the following error types were coded: 1) no response (i.e., response omission errors); 2) task substitutions (e.g., replacement of a verbal response with an action or vice-versa); 3) loss of content (e.g., acknowledgment that a response is required to a cue, but failure to recall the content); and (4) loss of time (i.e., performance of an intention greater than  $\pm 15\%$  before or after the target cue). If the participant makes no response to the PM cue, those are coded as “no response” errors and are presumed to be directly due to failure of PM (i.e., cue detection). Task substitution errors (e.g., intrusions and perseverations) are likely multi-determined, but presumed to be due to executive control deficits (e.g., Carey et al., 2004). Loss of content errors most likely reflect RM failures and loss of time errors seem to be due to difficulty with strategic monitoring or timing.

The Comprehensive Assessment of Prospective Memory (CAPM)

The CAPM is a self-report questionnaire used to assess PM. It takes approximately 10–15 minutes to complete. In this study we used only Section A containing 39 items relating to

frequency of PM failure in the last month. Items are rated on a five-point scale. This scale indicates that 1="never", 2 = "rarely", 3 = "occasionally", 4 = "often", and 5 = "very often".

Most items in Section A can be categorized into one of two subscales, IADL and BADL, established by Waugh (1999) using a principal component analysis. There are 23 items relating to IADL, such as "Leaving the iron on" and "Not remembering to pay bills". For the BADL subscale there are 10 items such as "Not locking the door when leaving home" and "Leaving water taps on." Given the "not applicable" category allowed, total scores and subscales scores were not used. Instead, for each participant three scores were calculated (total CAPM, IADL subscale, and BADL subscale) by summing the participant's ratings on the 1–5 scale for all completed items, and dividing by the total number of items less the not applicable items. Therefore, the possible range for mean total and subscale CAPM scores was 1–5, with higher scores indicating more frequently perceived PM failure.

#### Laboratory PM Task (Pereira et al., 2012)

The experimental session involved a practice phase for the ongoing task, followed by instructions for the PM task, a filled delay period and the main ongoing plus PM task. The effects of task variables on PM performance were examined on two measures: performance accuracy and latency for responding to the PM cues. Additionally, performance accuracy on the ongoing task, and response latency to non-PM cue items were recorded.

A simple computer-based activity in which participants had to sort 20 different noun words into two different categories (natural vs manmade) was prepared for the practice phase. For the PM cue-action pairings, two lists of 6 noun-verb pairs were compiled: one list comprised 6 related noun-verb pairs and the other 6 unrelated pairs. For the related list, noun-action words that had a mild semantic association ( $FSG < 0.1$ ; Nelson, McEvoy, & Schreiber, 1998) were

selected. In the unrelated list, the nouns from the related list were re-assembled with the verbs to create new pairs with no obvious associative relation between them. The word pairs had normative medium values of familiarity (Range = 3,71 - 4,59 on a scale of 1 to 7) and memorability (Range = 3,71 to 3,34 on a scale of 1 to 7); Molander & Arar (1998). For the main ongoing task, a set of 100 nouns (94 new and 6 cue-words) was created.

Participants were tested individually. They were informed that the session started with a practice task involving a simple computer-based activity in which they would have to allocate 20 different words into one of two different categories: natural or manmade by pressing the appropriate allocated key. Items remained on screen until the participant produced a response. This was followed by instructions for the PM task. Participants were presented with a set of 6 cue-action word pairs to learn. Half of the participants were presented 6 semantically mildly associated cue-action pairs. The other half of the participants was presented 6 cue-words that were not semantically associated in any obvious way. The mildly associated cue-action word pairs were as follows: ball – throw; coat – hang; flower – smell; lemon – squeeze; needle – prick; pencil – sharpen. All participants were informed that they would later perform a word-sorting task similar to the one performed during the practice phase. They were told that they would see a fixation in the centre of the computer screen for 3 seconds and that this would be followed by a sequence of words presented one at a time. They were asked to report, by pressing the relevant computer key, if the words belonged to the category “man-made”, or to the category “natural”. They were also told that some of the words of the task would have been presented in the previously encoded (prospective) set while others would be new. More specifically, participants were told that upon seeing the previously presented words (PM cue) they would have to press the computer key with the purple dot on it. After this, they would have to continue the word-sorting

task by pressing the appropriate key determining if that would be a natural or man-made stimulus.

Following presentation of all 6 items, participants were asked to complete a computerized version of the Stroop color-word interference task. This task required them to press a key on a response pad with a sticker corresponding to the color of the ink of the word displayed on the screen (red, yellow, blue or green). This lasted approximately 5 minutes.

Instructions for the main word-sorting (ongoing) task were then provided. However, none of the previously given information for the prospective task was given at that moment, if participants asked for more information about these previously given instructions at this point, they were simply told to do as they thought best.

Participants were reminded that they would see a fixation in the center of the computer screen for 3 seconds and that this would be followed by a sequence of words. They were asked to decide, by pressing the relevant computer key, if the words belonged to the category “man-made” (the computer key “z” had a sticker that said “manmade”) or to the category “natural” (the computer key “/” had a sticker that said “natural”). The 100 words (96 new, 6 PM cues) of the word-sorting task were then presented. Items remained on screen until the participant produced a response that he/she considered to be appropriate, by pressing a computer key. In the set of 100 words the cues were presented in the 8th, 20th, 44th, 55th, 82nd, and 99th position. These positions were not completely randomly selected because it was necessary to ensure that they were relatively evenly spread across the set of 100 words in such a way that the participant could not easily anticipate the exact position in which the cue would appear.



On completion of the word-sorting task, participants were asked if they remembered the instructions that had been given to them. This included describing what they thought they had been asked to do and also to recall all the 6 cue-action word pairs. As Maylor, et al. (2002) recommended, while studying prospective remembering it is crucial to separate PM failures from RM ones. This distinction might be particularly critical for studies of normal aging or dementia. Participants may perform poorly in a PM task not necessarily as a result of a PM failure, but because of a RM failure. Therefore, in the present study, participants were questioned at the end of the testing to ensure that they had successfully retained the PM instructions.

### Procedure

All participants were tested individually after obtaining informed consent. Total testing time was approximately 2 hours. Breaks were given if requested or if the participant complained of fatigue. The order of administration of the three tests was counterbalanced across participants.

### Data Analysis

None of the MIST measures were normally distributed as revealed by a series of Shapiro-Wilk  $W$  tests (all  $ps < .01$ ). Thus a series of non-parametric tests and Partial Eta Squared or Cohen's  $d$  (Hedges and Olkin, 1985) values were used to compare healthy adult and TBI samples on variables from the MIST. In all cases reported ANOVA results on raw data were confirmed with rank-transformed data; in no case was there a discrepancy in findings. Alpha was set at .05 and all analyses were two-tailed.

## Results

### MIST performance

Descriptive data are presented in Table 3. A repeated measures ANOVA for time delay (2 minute, 15 minute) x group (HA, TBI) revealed a significant main effect for time delay ( $F(1,143)=85.70, p<.001; \eta^2=0.39$ ), such that performance was superior at 2-min delays compared to 15-min delays; the main effect for group was also significant ( $F(1, 143)=44.26, p<.001; \eta^2=0.25$ ), such that healthy adults showed superior performance to those with brain injury; and the interaction was significant indicating that the effect of time was more pronounced for individuals with TBI ( $F(1, 143)=11.89, p<.01; \eta^2=.079$ ).

For type of cue (event, time) x group (HA, TBI), the main effect for type of cue was also significant ( $F(1,145)=89.36, p<.001; \eta^2=.40$ ), such that performance for event-based cues was superior to performance for time-based cues overall. The main effect for group was significant ( $F(1, 145)=53.40, p<.001; \eta^2=0.29$ ). The interaction was also significant indicating that the effect of cue was greater for individuals with TBI ( $F(1,145)=8.98; p<.01; \eta^2=.06$ ).

For type of response (action, verbal) x group (HA, TBI), the main effect of type of response was significant ( $F(1, 143)=17.61, p<.001; \eta^2=0.12$ ), such that performance on verbal response tasks was superior to that of action response tasks. The main effect of group was significant with the HA group performance superior to the TBI group ( $F(1,143)=44.60, p<.001; \eta^2=0.25$ ). The interaction was significant indicating that the effect of response was greater for the HA group ( $F(1, 145)=89.2, p<.01; \eta^2=.03$ ).

Individuals with TBI had significantly more PM (no response) errors, indicating no recall of the need to perform an intention ( $t(149)=10.21, p<.01; d^2=1.67$ ). They also performed significantly more poorly on the recognition items, indicating that RM is also impaired ( $t(149)$

=12.33,  $p < .01$ ;  $d^2 = 2.02$ ). On the more naturalistic 24-hour recall task, there was no difference between the groups. See Table 3 for the performance of the two groups on the MIST.

### CAPM Performance

For each participant, the average rating for items in both subscales (IADL and BADL) as well as for all 39 items (total CAPM) was calculated. The averaging of scores on the three scales was to take into account missing responses due to items being “not applicable”. Mean ratings and standard deviations for the items in the IADL and BADL subscales and total CAPM were calculated. On the CAPM, individuals with TBI indicated significantly lower BADL performance in daily life than controls ( $t(149) = 4.17, p < .05; d^2 = 0.68$ ), but there was no difference between the groups in terms of IADL or total performance. See Table 5.

### Laboratory PM Measure Performance

The laboratory PM measure data were analyzed using a 2 x 2 ANOVA of Group (HA, TBI) x Cue-Action relatedness (related, unrelated). There was a significant main effect of cue-action relatedness ( $F(1,141) = 19.20, p < .01; \eta^2 = 0.14$ ) such that PM performance was better overall when the cue was semantically related to the action. There was also a significant main effect of group, with HA individuals demonstrating significantly better performance than individuals with TBI ( $F(1,141) = 22.19, p < .01; \eta^2 = 0.18$ ). The interaction was not significant. See Table 4 for performance data on the two groups on this measure.

The proportion of cue-action word pairs that were remembered accurately at the end of the task was also calculated as a measure of retrospective recall. The number of correctly recalled cue-action word-pairs was calculated and there was no effect of cue-action relatedness

on the recall of the six word-pairs. However, recall was impaired in the group with TBI compared to the group of HA ( $t(149)=9.79, p<.01; d^2=1.60$ ). Comparison of Different Measures

Comparing measures, using Spearman rank-order correlations, for individuals with TBI, total correct on the laboratory task correlated significantly with a number of measures on the MIST. This included total correct and PM errors. See Table 6.

For the HA, the only significant correlation between the MIST and the laboratory measure was MIST recognition and total correct on the laboratory measure ( $r(98)=0.339; p<.05$ ). The laboratory task was not significantly correlated with the CAPM.

For the individuals with TBI, there were no significant correlations between the CAPM and either the MIST or the laboratory measure. For the HA the MIST total errors correlated significantly with BADL on the CAPM ( $r(98)=0.211; p<.05$ ).

When the TBI individuals were separated into low and high performance on the Stroop Word-Color Interference Test, using a median split, the groups differed in false positives on the laboratory measure, the MIST total correct, MIST recognition and PM errors but not the CAPM. See Table 7.

## **Discussion**

Overall, results of this study support previous research that has demonstrated that individuals with TBI are impaired compared to HA in PM performance and report significantly more PM deficits than HA (see Shum, Levin & Chan, 2011 for a review). More specifically, these results suggest that the impairment is most apparent when time delays exceed working memory span, self-initiated retrieval is required and the response requires an action. There is also a loss of RM content, most likely due to an independent RM impairment as seen on the

recognition tasks, although an error analysis from the MIST suggests that the PM impairment is considerably more prevalent than the RM impairment. Both groups showed a superiority for recognition over recall but the individuals with TBI were impaired in recognition, thus suggesting a deficit in both PM and RM systems. This is consistent with previous studies that have demonstrated separate significant effects in individuals with TBI (Pavawalla et al, 2012; Raskin et al, 2011).

The lack of a deficit in the TBI group when the delay period was short suggests that these data are more consistent with the multi-process theory than the PAM theory. In other words, under some conditions, individuals with TBI are able to perform adequately despite overall PM deficits.

The finding of a better performance on event-based as compared to time-based tasks is consistent with previous findings in populations with neurologic impairments (e.g., Raskin et al., 2011; Raskin et al., 2012). This has generally been explained by assuming that time-based tasks require more self-initiated retrieval and therefore greater cognitive resources allocation (e.g., Kvavilashvili, Cockburn & Kornbort, 2012), although some research has suggested that this may have to do with the extent to which the ongoing task and PM task share the same central executive resources (d'Ydewalle, Bouckart, & Brunfaut, 2001).

Results from the laboratory measure suggest that some of the difficulty with PM performance may be due to responding to an incorrect cue. Although no options for this are included in the MIST, the relationship of low cognitive control (i.e., poor Stroop performance) with PM errors may suggest a similar underlying pathology. There is some evidence to suggest that one aspect of PM performance (such as recognition of the PM cue) involves lateral prefrontal cortex plus precuneus (Burgess, Quayle, Frith, 2001) which may be the same regions

that mediate top-down attentional processes used for the Stroop (Banich, Milham, Atchley, Cohen, Webb, Wszalek, Kramer, Liang, Wright, Shenker, & Magin, 2000). This further lends support to the multi-process theory.

Results from the questionnaire suggest that the individuals with TBI perceived and reported greater difficulty with BADL activities than IADL activities, perhaps because failures at this level more significantly impact independence and generally represent a greater level of impairment (e.g., Harris, Jette, Campion, Cleary, 1986; Mitchell, Miller, Woodard, Davey, Martin, Burgess, Poon, 2011). This suggests that rehabilitation efforts might start with BADL activities that are perceived as more likely to be impaired.

These results support previous findings that the similarity of the retrieval cue and the intended action can influence the likelihood of successful PM performance. We found this to be true with the laboratory measure with the healthy adults, as has been shown previously (Ellis, 1996; McDaniel, Guynn, Einstein, & Breneisser, 2004; Pereira et al., 2012). We do not know of a previous study that has shown this same effect in individuals with TBI. This may point to a potential rehabilitation strategy for individuals with TBI in which individuals are trained to identify cues that are related to the intended action. In addition, this suggests that tests like the MIST, in which all event cues have highly related cue-action pairs, may be measuring only one aspect of performance on event-based cues.

When the groups are separated by adequate versus poor performance on a task of executive functioning, those with low executive functioning demonstrate significantly impaired performance on the clinical measure and the false positive response rate of the laboratory measure. Previous studies have not demonstrated a relationship between Stroop performance and PM in children with TBI (Ward, Shum, McKinlay, Baker & Wallace, 2007). Time-based

PM has been shown to be related to Stroop performance in adults with TBI (Mioni et al., 2012) but these data suggest that both time and event-based PM rely on these executive abilities and are specifically related to PM errors. Rule monitoring was shown to be related to PM performance on the Rivermead Behavioral Memory task in individuals with TBI, which is similar to the MIST (Paxton & Chiaravalloti, 2014). In contrast, these particular executive abilities do not seem to be related to the kind of PM required for the laboratory task or for awareness of deficits in daily life. This suggests that this aspect of executive functioning (error detection) may be related to some but not all features of PM.

Thus, these results support findings of separate contributions of different prefrontal regions to different aspects of PM performance (Burgess et al., 2000). The Stroop task has been previously shown to cause activation of anterior cingulate and inferior prefrontal regions (Leung et al., 2000). Therefore, the MIST may be tapping more into these functions whereas the laboratory task may be more related to cue detection and monitoring. In addition, responses on the questionnaire were not impacted by executive functioning and thus these questionnaires may be tapping into metacognitive awareness that is mediated by other prefrontal structures. This study is unable to answer this question and future studies that include imaging data on lesion location are necessary.

Comparing the clinical measure to the laboratory measure, it is interesting that there is little relationship between these two forms of PM measurement for the HA but that there was a strong relationship for those with TBI. Perhaps the tasks, designed not to be too difficult for those with TBI, were too simple for the HA, although ceiling effects were not noted. For those with TBI, these data suggest that both tasks are tapping into similar processes. In addition, it was not surprising that the greatest relationship was with the event-based items on the clinical

measure. This highlights the problem with using laboratory measures that include only event-based items if one is interested in a comprehensive picture of PM performance. The clinical measure was also related to false positive performance on the laboratory measure, in this case the number of PM errors. This may suggest that time-based remembering is related to self-monitoring more than event-based PM.

It is still unclear from this study why the questionnaire did not demonstrate a greater relationship to the other tasks. Perhaps this finding may be explained by the fact that the questionnaire is the only task that includes items requiring self-initiated planning. On the other hand, this finding is consistent with previous studies that have suggested that the issue of self-awareness is another factor that should be examined in more detail (Fleming et al., 2009). To clarify which of these is the likely explanation, future studies could take advantage of the significant other version of the CAPM. If significant correlations are found between those CAPM responses and the clinical or laboratory performance of those with TBI, the absence of significant relationships in this study are most likely explained as self-awareness deficits. In addition, perhaps performance on the CAPM could be compared to a diary study of PM performance in daily life, as rated by a significant other.

There are several limitations to this study. The first is the limitations of the singular measures given. For example, the laboratory measure includes on event-based items. Further the limitations of the MIST include less precise control of participant input than the computerized laboratory measure. The MIST may also not be measuring the same types of PM problems as are measured in daily life by the CAPM. The questionnaire leaves open issues of poor self-report. In addition, there were relatively few individuals with TBI included and no imaging data was available to know the specific areas of damage to the brain thus any



suggestions about different regions of prefrontal cortex having different functions are purely speculative.

Overall these results are consistent with the multi-process theory of PM. They support previous findings of PM deficits in TBI, with time-based items being more impaired than event-based items. Further, the results suggest that different aspects of PM performance may be being measured by different assessment techniques. Clinical assessments should take this into account when interpreting the effects of PM deficits on daily life. Further study of the differences between these techniques could yield important theoretical information about the nature of PM as well.

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Table 1. Demographic information or the two groups

	Healthy Adult N=100	Traumatic Brain Injury N=50	Group Differences
Age in years – Mean (sd)	29.78 (11.27)	31.54 (16.56)	n.s.
Education years Mean (sd)	14.41 (1.30)	15.13 (2.43)	n.s.
Handedness	83R 17 L	39 R 11 L	n.s.
Sex	51 F 49 M	32 M 18 F	

Table 2. Injury information for the group with TBI

	Mean (standard deviation)
GCS at scene or ED	11.87 (3.88)
Post traumatic amnesia (days)	1.35 (6.45)
Loss of consciousness (days)	11.23 (19.66)
Time post TBI (months)	29.32 (12.45)

Table 3. Performance of the two groups on the MIST

	Healthy Adult N=100 Mean (sd)	Traumatic Brain Injury N=50 Mean (sd)	<i>t(df=148)</i>	<i>d</i> <sup>2</sup>
Two-minute delay	7.80 (0.75)	6.94 (1.60)	2.82	0.46
15-minute delay	6.90 (1.30)	5.00 (2.12)	12.21***	2.00
Event cues	7.75 (0.65)	6.71 (1.36)	12.65***	2.07
Time cues	6.99 (1.11)	5.24 (1.71)	6.94**	1.13
Action response	7.28 (0.97)	5.76 (1.95)	10.40**	1.70
Verbal response	7.49 (0.72)	6.41 (1.33)	3.17	0.51
Total correct	44.15 (4.43)	35.82 (9.50)	13.91***	2.28
PM errors	0.14 (0.54)	1.83 (0.98)	18.46***	3.02
Recognition	7.68 (0.65)	7.00 (1.03)	41.53***	6.80
24 hour recall	0.76 (0.93)	0.50 (0.82)	1.71	0.28

\*\**p*<.01; \*\*\**p*<.001

Table 4. Performance of the two groups on the laboratory measure

	Healthy Control N=100 Mean (sd)	Traumatic Brain Injury N=50 Mean (sd)	t(df=148)	d <sup>2</sup>
Correct (max 12)	6.42 (4.42)	4.07 (4.57)	3.64	0.60
False Positives	0.26 (0.54)	0.50 (1.14)	12.04***	1.97
Correct (related) Proportion correct	0.81 (0.55)	0.71 (0.35)	7.12*	1.18
Correct (unrelated) Proportion correct	0.52 (0.49)	0.42 (0.32)	10.87**	1.78
Retrospective Recall	7.77 (4.22)	6.21 (5.54)	8.29*	1.36

\*p<.05; \*\*p<.01; \*\*\*p<.001

Table 5. Performance of the two groups on the CAPM

	Healthy Control N=100 Mean (sd)	Traumatic Brain Injury N=50 Mean (sd)	<i>t(df=148)</i>	<i>d</i> <sup>2</sup>
Basic ADL	1.54 (0.36)	1.80 (0.23)	4.17*	0.68
Instrumental ADL	1.96 (0.48)	1.94 (0.51)	1.02	0.17
Total Score on CAPM	1.83 (0.41)	1.81 (0.45)	1.43	0.23

\**p*<.05

Table 6. Significant correlations between the MIST and the laboratory measures for the individuals with brain injury (r values)

	Lab Measure Correct	Lab Measure False Positives	Correct (related)	Retrospective Recall
MIST Time Cues	0.641**			
MIST Event Cues	0.564**	0.219*	0.345*	
MIST Total Correct	0.628**			
MIST Recognition	0.608**			0.331*
MIST PM Errors	-0.446**			
MIST Total Errors	-0.702**	0.442*		

\*p&lt;.05; \*\*p&lt;.01



Table 7. Comparison of the TBI group with low performance on the Stroop to the TBI group with high performance on the Stroop.

	Stroop < 20 errors	Stroop >20 errors	<i>t(df=48)</i>
MIST total correct	0.91 (0.88)	1.82 (1.08)	12.17***
MIST PM errors	0.00 (0.00)	0.43 (0.53)	7.84**
MIST recognition	7.83 (0.45)	6.91 (1.05)	9.50**
MIST 24 hour recall	0.42 (0.76)	0.00 (0.00)	17.41***
BADL	1.52 (0.32)	1.34 (0.34)	2.33
IADL	1.93 (0.44)	1.67 (0.49)	3.94
Total CAPM	1.66 (0.49)	1.83 (0.36)	1.61
Correct on laboratory task	7.51 (3.96)	5.92 (4.22)	1.33
False Positives on laboratory task	0.28 (0.52)	1.09 (1.30)	9.16**

\*\* $p < .01$ ; \*\*\* $p < .001$