Less efficient pattern separation may contribute to age-related spatial memory deficits

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SPATIAL MEMORY DECLINE IN AGING

In the United States, Alzheimer’s disease (AD) is the most common cause of dementia in older adults and accounts for 60–80% of dementia cases (Alzheimer’s Association, 2012). In the year 2012, an estimated 5.4 million Americans will be diagnosed with AD, however, this number is projected to increase to 11–16 million by 2050 (Alzheimer’s Association, 2012). As a result of the aging “baby boom” generation and increasing longevity in the US population, the disease is a growing public health concern with costs estimated to reach $200 billion in 2012. Although a number of risk factors for AD have been discussed (e.g., diagnosis of mild cognitive impairment (MCI), family history of AD, apolipoprotein epsilon E4 allele), one of the most well-documented risk factors for the disease is increasing age (Kamboh, 2004). Therefore, a major aim of recent research has focused on identifying early indicators of cognitive dysfunction in older adults.

A variety of cognitive functions may be affected in older adults, however, one of the most commonly reported deficits associated with aging is memory loss. Although not all memory domains are equally affected by aging (e.g., source vs. item memory), some domains such as spatial memory appear to be particularly sensitive to age-related change. Age-related spatial memory decline has been well-documented in both humans (reviewed by Iachini et al., 2009) and animal models (reviewed by Sharma et al., 2010). Spatial memory deficits also have been well-documented in older adults diagnosed with AD (Sahakian et al., 1988; Kessels et al., 2005) or mild cognitive impairment (MCI: Alescio-Lautier et al., 2007; deIpolyi et al., 2007; Hort et al., 2007). Some evidence suggests that spatial memory deficits even may serve as an early indicator of AD (Buccione et al., 2007; deIpolyi et al., 2007; Hort et al., 2007). Age-related spatial memory deficits may result from changes in a variety of brain regions including the hippocampus, temporal lobes, and the frontal-parietal network (Iachini et al., 2009). In particular, spatial memory decline associated with aging has been suggested to result from age-related changes in the hippocampus across species (reviewed by Barnes, 1988). However, it is only recently that studies have begun to examine how age-related changes in particular subregions of the hippocampus may affect specific mnemonic processes. One such process is spatial pattern separation, a mechanism that may be essential for the accurate formation and retrieval of spatial memories. Less efficient spatial pattern separation in older humans and animals may contribute to impairments in spatial memory, particularly in situations when spatial interference is high.

PATTERN SEPARATION

Pattern separation is a mechanism for separating partially overlapping patterns of activation so that one pattern may be retrieved as separate from other similar patterns. A pattern separation mechanism may be critical for reducing potential interference among similar memory representations to enhance memory accuracy (Gilbert and Brushfield, 2009). A number of models have suggested that the hippocampus supports pattern separation (Marr, 1971; McNaughton and Nadel, 1990; O’Reilly and McClelland, 1994; Shapiro and Olton, 1994; Rolls and Kesner, 2006; Kesner, 2007; Myers and Scharfman, 2009; Rolls, 2010). In particular, the dentate gyrus (DG) and CA3 subregions of the hippocampus have been shown to play a critical role in

Keywords: hippocampus, dentate gyrus, aging, spatial memory, interference, pattern separation

Spatial memory deficits have been well-documented in older adults and may serve as an early indicator of mild cognitive impairment (MCI) or Alzheimer’s disease (AD) in some individuals. Pattern separation is a critical mechanism for reducing potential interference among similar memory representations to enhance memory accuracy. A small but growing literature indicates that spatial pattern separation may become less efficient as a result of normal aging, possibly due to age-related changes in subregions of the hippocampus. This decreased efficiency in spatial pattern separation may be a critical processing deficit that could be a contributing factor to spatial memory deficits and episodic memory impairment associated with aging. The present paper will review recently published studies in humans, non-human primates, and rodents that have examined age-related changes in spatial pattern separation. The potential basic science, translational, and clinical implications from these studies are discussed to illustrate the need for future research to further examine the relationship between spatial pattern separation and brain changes associated with aging and neurodegenerative disease.
tern separation activity in the DG/CA3 regions of aged humans. A recent study by Yassa et al. (2011a) reported that reduced pattern completion at the expense of processing new information retrieved using partial cues. These age-related changes may cause that allows a complete representation of stored information to be compromised in a subset of older adults. In an attempt to replicate performance of both young and older adults increased as a function of increased age-related differences in maintaining a pattern separation-pattern completion equilibrium.

Given the well-documented role of the DG in supporting pattern separation and the potential susceptibility of this region to age-related change, recent behavioral studies have begun to examine pattern separation in older humans. Using a continuous recognition paradigm developed by Kirwan and Stark (2007), Toner et al. (2009) reported that non-demented older adults are impaired on a pattern separation task for visual object information. A subsequent fMRI study supported these findings and additionally demonstrated increased age-related activity in the DG and CA3 subregions when pattern separation demands were high on a visual object task (Yassa et al., 2010a).

**SPATIAL PATTERN SEPARATION IN OLDER HUMANS**

Recent studies also have begun to examine spatial pattern separation deficits in older adults. A study by Stark et al. (2010) reported that spatial pattern separation is impaired in a subset of older adults. Participants studied unique pairs of pictures and later were asked to identify whether the pictures were both in the same location as before or whether one of the pictures was in a different location. In the same condition, neither of the pictures in a pair was moved. In the three different conditions (close, medium, far) the location of one of the pictures in a pair was moved by varying both the distance and the angle from the original location. Older adults were separated into an aged impaired group and aged unimpaired group, based on standardized word learning task performance. Analyses revealed performance deficits on the close, medium, and far trials for the aged impaired group relative to both the aged unimpaired and young groups. Therefore, the findings suggest that spatial pattern separation may be impaired in a subset of older adults.

A subsequent study by Holden et al. (2012) examined the ability of cognitively normal young and older adults to perform a spatial memory task involving varying degrees of spatial interference to assess spatial pattern separation. During the sample phase of each trial, a circle appeared briefly on a computer screen. The participant was instructed to remember the location of the circle. During the choice phase, two circles were displayed simultaneously and the participant was asked to indicate which circle was in the same location as the sample phase circle. The two choice phase circles were separated by one of four possible spatial separations across trials: 0, 0.5, 1.0, and 1.5 cm. Smaller separations are likely to create increased overlap among memory representations, which may result in heightened interference and a greater need for pattern separation. Consistent with this hypothesis, the performance of both young and older adults increased as a function of increased spatial separation. However, young adults outperformed older adults, suggesting that spatial pattern separation may be less efficient in older adults. In an attempt to replicate the findings of Stark et al. (2010), the older adult group was separated into older impaired and older unimpaired groups based on the well-documented role of the DG in supporting pattern separation activity in the DG/CA3 regions of aged humans. A recent study by Yassa et al. (2011a) reported that reduced pattern completion at the expense of processing new information retrieved using partial cues. These age-related changes may cause that allows a complete representation of stored information to be compromised in a subset of older adults.
on performance on a standardized delayed word recall measure. Consistent with Stark et al. (2010), the older impaired group was significantly impaired relative to both the older unimpaired and young groups.

**Spatial Pattern Separation in Older Animals**

Kubo-Kawai and Kawai (2007) conducted a study in aged monkeys that is similar in nature to the two aforementioned studies conducted in older humans. This study tested young and older monkeys on a delay non-match-to-position task involving manipulations of the spatial distance between two locations. During the sample phase, a plate was positioned to cover a baited food well along a single row of four food wells. The animal displaced the plate to receive a reward. During the choice phase, an identical plate was positioned to cover the unbaited sample phase well (incorrect choice) and a second identical plate was positioned to cover a different baited food well in one of the three remaining locations (correct choice). The distance between the two choice phase plates was manipulated across trials and included separations of 12, 24, and 36 cm. The data showed that the ability of both young and older monkeys to choose the correct plate improved as a function of increased spatial separation between the two plates during the choice phase. However, no age-related differences in performance were detected across separations. In a second experiment, the sample phase followed the same procedure described for the previous experiment. However, on the choice phase, two rows of food wells were used. The plate covering the unbaited sample phase food well (incorrect choice) was located on one row, whereas the plate covering the new location was positioned to cover a baited food well (correct choice) on a different row of wells. Spatial separations of 15, 25.6, and 37.1 cm were used to separate the plates across choice phase trials. The data revealed that older monkeys were impaired relative to young monkeys across all three separations, with the largest group differences found on trials involving the closest 15 cm separation. One could speculate that smaller separations were likely to create increased overlap among memory representations, which may result in heightened interference and a greater need for pattern separation. Although this study was not designed or interpreted by the authors to assess pattern separation, these findings in aged monkeys are in accordance with the findings from the studies involving older humans.

A recent study from our lab (Gracian et al., unpublished observations) provides further behavioral evidence that spatial pattern separation may be impaired in aged animals. Young and old rats were trained on a radial eight-arm maze to discriminate between a rewarded arm and a non-rewarded arm that were either adjacent to one another (high spatial interference) or separated by a distance of two arm positions (low spatial interference). Performance on this task recently was shown to be dependent on the DG hippocampal subregion (Morris et al., 2012). Gracian et al. showed that young and old rats committed similar numbers of errors in the separated condition. However, in the adjacent condition, old rats committed significantly more errors compared to young rats. The results suggest that decreased spatial pattern separation in aged rats may impair performance on the adjacent condition, which involved greater spatial interference among distal cues. However, performance increased in the separated condition when there was less overlap among distal cues and less need for pattern separation.

A recently published study by Marrone et al. (2011) provides further neurobiological insight into how age-related changes in the DG may affect pattern separation and spatial memory. In this study, young and aged rats were allowed to explore either: (1) the same environment on two occasions or (2) one environment followed by a different environment. A marker of cellular activity (zif268/egr1) was used to examine granule cell activity during exploration of the environments. Older animals were found to recruit distinct populations of granule cells during exploration of the same environment on two different occasions, which the authors interpret to be an indication of greater pattern separation. However, when the aged animals visited different environments, the reliability of repeated zif268 expression in young and aged animals was comparable, thus the age-related increase in pattern separation was no longer apparent. The study also found that increased pattern separation in the similar contexts was correlated with a reduced ability of older animals to disambiguate similar contexts during a sequential spatial recognition task. The authors conclude that spatial memory performance in aged animals is most impaired in situations where interference is increased, presumably due to decreased pattern separation. Collectively, the aforementioned studies offer evidence that spatial pattern separation may become less efficient as a result of aging.

**Implications**

Age-related memory decline has been suggested to stem from subregion-specific epigenetic and transcriptional changes in the hippocampus (Penner et al., 2010). For example, neurogenesis is reduced in aged animals (Kuhn et al., 1996) and is related to decreased hippocampal volume and impaired performance on hippocampal-dependent tasks (Driscoll et al., 2006). These newborn neurons may be involved in mnemonic processes particularly dependent on the DG subregion, such as pattern separation (Clelland et al., 2009; Aimone et al., 2010, 2011; Creer et al., 2010; Deng et al., 2010; Sahay et al., 2011), whereas older DG cells may contribute to pattern completion (Nakashiba et al., 2012). Sahay et al. (2011) suggest that interventions that increase neurogenesis during adulthood may have clinical implications for reversing age-related impairments in pattern separation and associated DG dysfunction. Therefore, the development of behavioral tasks sensitive to age-related changes in spatial pattern separation may have implications for future studies of neurogenesis.

Age-related changes in spatial pattern separation also may contribute to episodic memory deficits, which have been well-documented in older adults (Rand-Giovannetti et al., 2006) and are a prominent feature of AD that may be detectable several years before disease onset (Bondi et al., 1999). Episodic memory also is impaired in non-demented older adults who are at risk for AD based on genetics (Saunders et al., 1993) or a diagnosis of MCI (Hodges et al., 2006). One key feature of episodic memory that differentiates it from other types of memory is that the elements of an episodic memory must be associated into a context to demarcate the episode in space and time. In addition, a pattern separation mechanism may be necessary to separate
the elements of different episodic memories to avoid interference (Gilbert et al., 2001). It is possible that less efficient spatial pattern separation in older adults may result in poorer memory due to increased interference among the spatial components of episodic memories. The identification of a key mnemonic processing deficit in pattern separation may result in behavioral interventions that structure daily living tasks to mitigate interference in the spatial domain and potentially improve spatial and episodic memory in older adults.

Normal and pathological aging may have differential effects on hippocampal subregions. The DG subregion may be particularly susceptible to age-related changes in humans, however, there may be less impact on pyramidal cells in the CA subregions (Small et al., 2002). In contrast, the CA subregions may be more vulnerable to pathological changes associated with AD (Braak and Braak, 1996; West et al., 2000; Price et al., 2001; Apostolova et al., 2010). As mentioned previously, a primary goal in AD research is to identify risk factors and preclinical markers of the disease in older adults. Given the differential effects of normal aging and AD on the various subregions of the hippocampus, tasks that are sensitive to dysfunction in particular subregions, such as measures of pattern separation, may help to differentiate between cognitive impairment associated with normal aging and pathological changes associated with AD. In support of this idea, a recent study by Yassa et al. (2010b) reported that individuals with amnestic MCI show impairments compared to healthy older adults on an object recognition memory task that taxes pattern separation. In addition, structural and functional changes were observed in the DG/CA3 hippocampal subregions of these individuals.

CONCLUSIONS

In conclusion, spatial memory deficits have been well-documented in older adults and may serve as an early indicator of MCI or AD in some individuals. The operation of a pattern separation mechanism may be critical for reducing interference among similar memory representations to enhance memory accuracy. Evidence suggests that brain regions critical to pattern separation, including the DG and CA3 hippocampal subregions, may be particularly susceptible to adverse age-related changes. A small but growing literature indicates that spatial pattern separation may become less efficient as a result of normal aging. This decreased efficiency in pattern separation may contribute to spatial memory deficits and episodic memory impairment associated with aging. Based on the aforementioned studies, it is clear that additional research is needed to examine the relationship between spatial pattern separation and brain changes associated with aging and neurodegenerative disease.

ACKNOWLEDGMENTS

A portion of the research reviewed in the paper was supported by a National Institutes of Health Grant (#AG034202) from the National Institute on Aging awarded to Paul E. Gilbert.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Received: 29 March 2012; paper pending published: 27 April 2012; accepted: 14 May 2012; published online: 29 May 2012.

Citation: Holden HM and Gilbert PE (2012) Less efficient pattern separation may contribute to age-related spatial memory deficits. [Front. Ag. Neurosci. 4:9. doi: 10.3389/fnagi.2012.00009]

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