



# The Shifting Architecture of Cognition and Brain Function in Older Adulthood

Perspectives on Psychological Science 2019, Vol. 14(4) 523–542 © The Author(s) 2019
Article reuse guidelines: sagepub.com/journals-permissions
DOI: 10.1177/1745691619827511
www.psychologicalscience.org/PPS

\$SAGE

R. Nathan Spreng<sup>1</sup> and Gary R. Turner<sup>2</sup>

<sup>1</sup>Department of Neurology and Neurosurgery, McGill University, and <sup>2</sup>Department of Psychology, York University

#### **Abstract**

Cognitive aging is often described in the context of loss or decline. Emerging research suggests that the story is more complex, with older adults showing both losses and gains in cognitive ability. With increasing age, declines in controlled, or fluid, cognition occur in the context of gains in crystallized knowledge of oneself and the world. This inversion in cognitive capacities, from greater reliance on fluid abilities in young adulthood to increasingly crystallized or semanticized cognition in older adulthood, has profound implications for cognitive and real-world functioning in later life. The shift in cognitive architecture parallels changes in the functional network architecture of the brain. Observations of greater functional connectivity between lateral prefrontal brain regions, implicated in cognitive control, and the default network, implicated in memory and semantic processing, led us to propose the *default-executive coupling hypothesis of aging*. In this review we provide evidence that these changes in the functional architecture of the brain serve as a neural mechanism underlying the shifting cognitive architecture from younger to older adulthood. We incorporate findings spanning cognitive aging and cognitive neuroscience to present an integrative model of cognitive and brain aging, describing its antecedents, determinants, and implications for real-world functioning.

#### **Keywords**

representation, default-mode network, semantic knowledge, executive

Across society, from public policy to popular culture, cognitive aging has become synonymous with cognitive decline. This unfortunate heuristic is inaccurate in the context of a growing body of evidence showing that the full spectrum of cognitive aging encompasses both gains and losses (Park, Polk, Mikels, Taylor, & Marshuetz, 2001; Fig. 1). In this review, we characterize the evidence for a shift in the architecture of cognition across the life span, with the hope of better characterizing the full spectrum of cognitive aging. We then describe how the changing architecture of brain function provides a striking parallel to these cognitive changes. We conclude by offering a novel model of neurocognitive aging to better bridge our understanding of cognitive and brain aging, to generate predictions for both gains and losses, and finally to propose a putative neural mechanism connecting these trajectories of cognitive and brain aging.

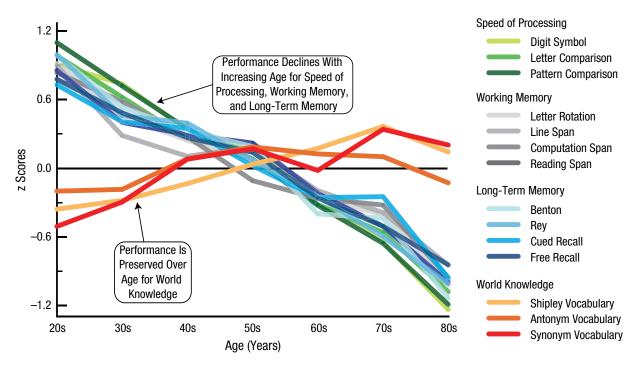
### **Dual Intelligences and the Semanticization of Cognition**

An accurate portrayal of aging cognition must consider the dual nature of fluid and crystallized intellectual ability, and their respective trajectories, across the life span. Fluid intelligence indexes cognitive operations that require goal-directed, or controlled, processing. We refer to these fluid abilities throughout the review as cognitive-control processes, necessary for directing attentional resources to relevant aspects of the environment, inhibiting distractions, and flexibly allocating processing resources to sustain goal-directed behaviors (Carpenter, Just, & Shell, 1990; Cattell, 1971). Crystallized intelligence indexes the collective store of semantic knowledge about ourselves and the world that is accumulated over the life course (Cattell, 1971). We refer to these crystallized abilities here as *semantics*, encompassing a vast repertoire of prior knowledge, including routines, habits, stereotypes, and schemas, as well as the thoughts and behaviors, one repeatedly

#### **Corresponding Author:**

R. Nathan Spreng, Laboratory of Brain and Cognition, Montreal Neurological Institute, Department of Neurology and Neurosurgery, McGill University, 3801 University St., Montreal, Quebec H3A 2B4, Canada

E-mail: nathan.spreng@gmail.com



**Fig. 1.** The shifting architecture of cognition across the adult life span. Cool colors represent age-related changes on tasks that have greater reliance on cognitive control and speeded processing. Warm colors represent age-related changes on tasks that have a greater reliance on semantics or prior-knowledge representations. As can be seen, the architecture of cognition undergoes a striking shift across the adult life span, a process we refer to as the *semanticization of cognition*. Republished with permission of Annual Reviews, Inc., from *The Adaptive Brain: Aging and Neurocognitive Scaffolding*, by Park, D. C., & Reuter-Lorenz, P., in *Annual Review of Psychology*, Vol. 60, Copyright © 2009; permission conveyed through Copyright Clearance Center, Inc.

encounters over a lifetime (Jefferies, 2013; Ralph, Jefferies, Patterson, & Rogers, 2017).

These two facets of cognition, control processes and crystallized knowledge, age differently. Cognitivecontrol abilities show marked and near linear declines with increasing age (Park et al., 2001; Verhaeghen & Cerella, 2002). In contrast, semantics continue to accumulate and are preserved well into later life (Park et al., 2001; Verhaeghen, 2003). Although these dual trajectories have been thoroughly described as independent facets of cognitive aging, surprisingly little research has examined how they interact. This is rather remarkable given the striking inversion of cognitive abilities that occurs across the adult life course (Park et al., 2001; see also Fig. 1). We refer to this shift as the semanticization of cognition in older adulthood, and the impacts can be seen across a diverse array of cognitive domains from memory (e.g., Umanath & Marsh, 2014) to socioemotional functioning (e.g., Carstensen, Fung, & Charles, 2003), as well as in multidomain abilities such as decision making and problem solving, with implications for real-world functioning and independence (e.g., Agarwal, Driscoll, Gabaix, & Laibson, 2009; Y. Li, Baldassi, Johnson, & Weber, 2013). Yet how and why this shift occurs has not been fully explored.

### **Models of Neurocognitive Aging**

Two of the most commonly reported patterns of functional brain change in older adulthood are (a) greater recruitment of prefrontal cortical regions, implicated in cognitive control, and (b) reduced suppression of the default network, an assembly of functionally connected brain regions implicated in one's ability to access mnemonic representations of oneself and the world (Andrews-Hanna, Smallwood, & Spreng, 2014). Increased recruitment of lateral prefrontal brain regions has been described as hemispheric asymmetry reductions in old age (HAROLD; Cabeza, 2002) or a posterior-to-anterior shift in aging (PASA; Davis, Denis, Deselaar, Fleck, & Cabeza, 2008). These early influential models of neurocognitive aging emerged from an abundance of evidence showing that older adults display a consistent pattern of enhanced prefrontal recruitment, including greater bilateral activation, across many cognitive tasks. Enhanced recruitment is attributed to greater demands for cognitive-control processes, necessary in the context of degraded signaling from earlier processing zones located in posterior brain regions (Li, Lindenberger, & Sikstrom, 2001; S. C. Li & Rieckmann, 2014; Payer et al., 2006). Later, models emerged to suggest that this dedifferentiated

pattern of enhanced prefrontal recruitment serves to support cognitive functioning in the context of structural and functional changes in later life. This pattern of functional brain adaptation has been characterized in the compensatory recruitment of neural circuits hypothesis (CRUNCH; Reuter-Lorenz & Cappell, 2008) or the scaffolding theory of aging cognition (STAC; Park & Reuter-Lorenz, 2009).

Neurocognitive aging has also been explored through the lens of large-scale functional brain networks (Damoiseaux, 2017). The default network, a collection of intrinsically connected brain regions, including the posterior cingulate cortex, medial prefrontal cortex, inferior parietal lobule, and medial and lateral temporal lobes, has been particularly implicated (Hafkemeijer, van der Grond, & Rombouts, 2012). This network is typically activated during internally directed cognitive processes that rely on access to stored mnemonic representations to guide goal-directed behaviors (Andrews-Hanna et al., 2014). In contrast, default-network brain regions are suppressed during the performance of externally directed tasks (Buckner, Andrews-Hanna, & Schacter, 2008). Age-related changes in the default network include reduced suppression, as well as decreased within-network and increased between-network connectivity, each of which is poorly modulated by task context (e.g., Rieck, Rodrigue, Boylan, & Kennedy, 2017; Spreng & Schacter, 2012; for reviews, see also Damoiseaux, 2017; Hafkemeijer et al., 2012).

# Connecting Cognitive and Functional Brain Change: A New Proposal

Considered together, these functional brain changes, involving regions implicated in cognitive control (lateral prefrontal cortex) and access to prior-knowledge representations (default network), closely parallel the shift in cognitive architecture described earlier. There has been little research, however, to connect these patterns of cognitive and brain aging. Our work provided some of the first evidence that lateral prefrontal engagement and reduced default-network suppression are functionally coupled in older adulthood. This led us to propose the default-executive coupling hypothesis of aging (DECHA; Turner & Spreng, 2015). This hypothesis predicts that as goal-directed cognition becomes less dependent on declining control resources and increasingly influenced by prior knowledge (i.e., the semanticization of cognition), the default network is engaged and becomes increasingly—and inflexibly—coupled with lateral prefrontal brain regions (Fig. 2).

Although broadly consistent with other neurocognitive-aging theories in positing a role for increased prefrontal engagement, DECHA differs from these earlier compensatory accounts by highlighting the role of cognitive context in determining whether these brain changes are functionally adaptive or maladaptive. Put simply, if access to semantic knowledge is relevant to the current task, greater default-executive coupling should be adaptive (Adnan, Beaty, Silva, Spreng, & Turner, 2019; Spreng et al., 2014). In contrast, if prior knowledge is irrelevant or distracting, this pattern should lead to poorer performance (Rieck et al., 2017; Turner & Spreng, 2015). DECHA reflects a growing body of evidence showing that this coupling pattern is increased, and poorly modulated by task context, in older adulthood (Rieck et al., 2017; Spreng & Schacter, 2012). Here we suggest that DECHA opens a novel avenue for exploring these parallels in cognitive and brain aging. Critically, as a putative neural mechanism, reflecting both declining fluid abilities and increasing semantic knowledge (i.e., the semanticization of cognition), DECHA can characterize a broad spectrum of aging cognition, encompassing both gains and losses in adaptive, goal-directed behavior in later life.

### **Objectives**

In the sections that follow, we expand on the findings introduced thus far to provide a comprehensive review of the cognitive and functional brain changes that occur with age, how they promote the semanticization of cognition in later life, and how this shift in cognitive architecture parallels the shift in brain function described in DECHA. We begin by describing the shifting architecture of cognition from younger to older adulthood and highlight the impacts, both negative and positive, on real-world functioning. We then directly interrogate our hypothesis that the semanticization of cognition is attributable to the interaction of increased semantics and declining control. We argue that this shifting architecture leads to an enduring shift in cognitive mode, from exploration and novelty seeking in young adulthood to greater exploitation of existing knowledge stores in later life. We then turn to brain aging and provide a survey of the mounting evidence for enhanced recruitment of lateral prefrontal brain regions and reduced modulation of the default network across the adult life span, in addition to evidence for the functional coupling of these regions with advancing age. In the final section we bring together the evidence supporting our novel, cross-disciplinary theoretical framework, DECHA, and argue that it represents an integrative neural mechanism underlying the shifting architecture of cognition in older adulthood.

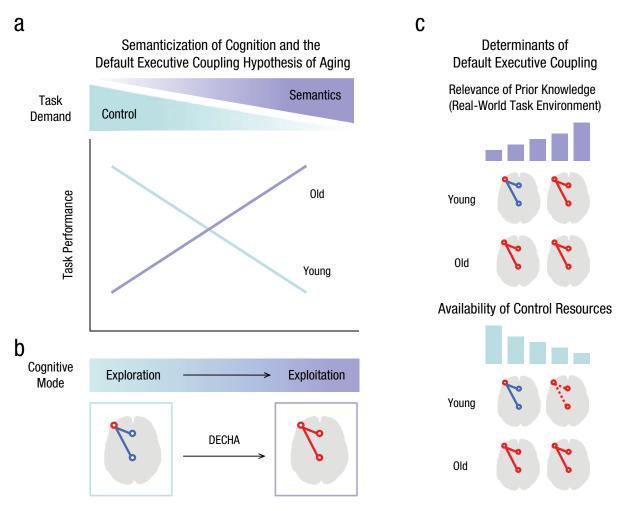


Fig. 2. Integrated model of cognitive and brain aging. Shifting task demands, from controlled processing to greater reliance on semantics (prior knowledge), affect task performance differently for younger and older adults (a). When control demands are high, there is a relative performance benefit for younger adults (left side of the graph). As prior knowledge becomes more goal-relevant, the performance advantage shifts to older adults (right side of the graph). Greater reliance on semantic knowledge representations, in the context of declining control resources (see Fig. 1), results in a shift in cognitive mode from exploration to exploitation in later life (b). Schematics of the brain represent the default-executive hypothesis of aging (DECHA). The bottom-left brain schematic reflects greater reliance on cognitive-control processes or the exploratory cognitive mode. This mode is associated with the engagement of lateral prefrontal brain regions (red circle) and suppression of default-network brain regions (blue circles). Blue lines indicate negative functional connectivity between default and executive regions (i.e., greater frontal activation in the context of default-network suppression). The bottom-right brain schematic reflects greater reliance on prior knowledge or an exploitative cognitive mode. This mode is associated with increased and positive functional connectivity between lateral prefrontal regions and the default network (red lines). We hypothesize that this pattern of increased functional connectivity serves as a neural-network mechanism promoting the semanticization of cognition in later life. As the relevance of prior knowledge increases (c, top) or cognitive control declines (c, bottom), functional connectivity between prefrontal and default network brain regions flexibly adapts in young people (shift from blue to red lines). In contrast, default-executive coupling is less flexible in older adults and is poorly modulated by task context. (Note that direct evidence for frontal-default coupling in young people when control resources are reduced is lacking in the literature; therefore, connectivity patterns are represented by dotted red lines.) Adapted with permission of the American Psychological Association, from Reward Processing and Risky Decision Making in the Aging Brain, by Samanez-Larkin, G. R., & Knutson, A., in The Neuroscience of Risky Decision Making (V. Reyna & V. Zayas, Eds.), Copyright © 2014; permission conveyed through Copyright Clearance Center, Inc.

## The Shifting Cognitive Architecture of Aging

In this section we review evidence for the semanticization of cognition in older adulthood and how this manifests itself within specific domains, such as memory or social cognition, as well as in multidomain abilities such as planning, problem solving, and decision making. Our intention is not to provide a comprehensive survey of the impact of prior knowledge on cognitive functioning across the life span. Excellent reviews of this topic can be found elsewhere (e.g., Umanath & Marsh, 2014).

Our goal is to provide an overview of the most pertinent research in this area as a basis for appreciating the parallels between cognitive and brain aging we draw later in the article.

### Cognitive aging and semanticknowledge gains

The semanticization of cognition has been characterized as an age-related transition toward reliance on prior knowledge, including semantics, conceptual-level knowledge, and schematic representations. This occurs in the context of declining access to detailed representations of the past (e.g., episodic memories), as the unique and arbitrary associations that form the basis of these discrete representations decay over time (Craik & Bialystok, 2006). There have been numerous, often competing, models put forth to explain age-related cognitive decline. Although a broader review of neurocognitive aging models is beyond the scope of this article (see Anderson & Craik, 2017; Park & Reuter-Lorenz, 2009), age-related declines in inhibitory capacity (Hasher & Zacks, 1988) and reduced speed of processing (Salthouse, 1996) are among the most often cited accounts. Evidence, however, is emerging to suggest that the accrual of semantic knowledge over the life course may be a superordinate factor, interacting with specific declines in control processes secondary to frontal brain changes (e.g., inhibition) or slowed speed of processing, to accelerate age-related cognitive changes. For example, a larger store of semantic representations, in the context of reduced inhibitory capacity, would be more frequently activated—automatically or inadvertently across a broader range of cognitive contexts (Hoffman, 2018). Likewise, an increased store of semantics would require longer searches through an expanded solution space, slowing processing speeds in later life (Blanco et al., 2016; Ramscar, Hendrix, Shaoul, Milin, & Baayen, 2014). Next we explore the impacts of the semanticization of cognition first with respect to domain-specific impact (verbal functioning, memory, associative learning, cognitive control) and then followed by multidomain and more real-world implications (problem solving, decision making).

### Domain-specific impacts

Beyond interactions with specific control processes, the impact of an age-related shift toward semanticized cognition has been investigated across multiple cognitive domains. With respect to verbal ability, older adults demonstrate superior performance on tasks assessing vocabulary size (Park et al., 2001; Verhaeghen, 2003). Older-adult recognition for previously studied words is

also better for those embedded within sentences than for those presented as stand-alone items. This has been attributed to linguistic expertise honed over a lifetime of reading (Matzen & Benjamin, 2013). Further, older, but not younger, adult vocabulary scores can significantly predict memory performance on a cued recall test (Hedden, Lautenschlager, & Park, 2005).

In contrast, older adults have greater difficulty learning alternate versions of fairy tales, presumably because the overlearned, standard narratives proactively interfere with the encoding of the nonstandard versions (Dalla Barba, Attali, & La Corte, 2010). A similar effect has been observed during the learning of misspelled words (MacKay, Abrams, & Pedroza, 1999) as well as incorrect mathematical solutions (Ruch, 1934). In each of these cases, prior knowledge disrupts learning and subsequent task performance.

However, the influence of prior knowledge may also protect against cognitive distortions. As reviewed in Umanath and Marsh (2014), an interesting example of this phenomenon involves the Moses illusion, in which participants are required to answer both factually correct and incorrect questions (e.g., "How many animals of each species did Moses lead to the ark?"). Although older adults failed to notice factual errors at the time of the study (Noah is the protagonist of the ark narrative), they were less likely than younger adults to provide the incorrect responses on a subsequent test (e.g., "Which biblical figure led pairs of animals to an ark?"). The older-adult advantage was observed even when older and younger participants were matched on prior knowledge (Umanath & Marsh, 2012).

This effect has also been observed in more ecologically valid contexts. Older adults are less likely than younger adults to remember invalid versus valid prices for grocery items (Amer, Giovanello, Grady, & Hasher, 2018; Castel, 2005). They are also less likely to exhibit susceptibility to the illusory-truth phenomenon, which describes the tendency of younger adults to perceive facts repeated more often as true. Older adults can draw from their larger store of semantic knowledge to avoid this illusion, using prior knowledge to circumvent the learning of false information as a function of exposure (Brashier, Umanath, Cabeza, & Marsh, 2017). In general, differences in memory performance between younger and older adults are reduced or eliminated when tobe-remembered information is consistent with prior knowledge. In their review, Umanath and Marsh (2014) put it simply: "The more prior knowledge older adults can apply, the smaller the age difference in memory" (p. 214).

Although the semanticization of cognition can protect older adults from encoding false or distorting information when it is incongruent with one's prior knowledge,

this same process can also lead to memory distortions when prior knowledge can shape or influence encoding processes. An example of this is the phenomenon referred to as false-memory formation. Here, reliance on prior knowledge leaves older adults more susceptible to the formation of imprecise or incorrect memories, or illusory associations, during encoding and retrieval of conceptually linked word lists (Schacter, Koutstaal, & Norman, 1997). Using the Deese-Roediger-McDermott (DRM) paradigm (Deese, 1959; Roediger & McDermott, 1995), participants studied word lists that share a common underlying theme, for example, words associated with cold (frost, snow, ice, etc.). On a subsequent recognition test, older adults more frequently endorsed target lures, words related to the theme but were not on the original study list (e.g., frozen). The authors argued that these "false memories" are generated as the result of richer associations for study items with a larger store of semantic representations in older versus younger adults. In a separate experiment and in line with this idea, Koutstaal et al. (2003) implemented a visual adaptation of the DRM paradigm in which participants studied a series of abstract images and were asked to identify the previously studied items from among a series of new images that included visually similar items on a later recognition probe. The older participants in the study, unlike those from Roediger and McDermott (1995), did not endorse more visually similar lures than the younger participants. However, in a separate condition, concrete labels were assigned to the images at the study phase (e.g., watch, truck, lamp, bread). In this condition, older adults did endorse more target lures than younger adults, providing strong evidence that the formation of false memories was related to increased semantic associations as a result of the verbal labels assigned to the

Semanticization of memory has also been observed during the recollection of personal past events or autobiographical recall. During the Autobiographical Interview, a structured interview to assess recollections of one's personal past (Levine, Svoboda, Hay, Winocur, & Moscovitch, 2002), older adults report proportionately more personal semantic information (e.g., "I loved playing basketball in high school") than episodic details of their past (e.g., "I shot the winning basket in the final game of the season in my junior year"). As we describe later in the review, our work is beginning to show that the semanticization of personal past memories may be associated with changes in the network architecture of the brain in older adults (Spreng et al., 2018).

The influence of prior knowledge on cognition in older adulthood also extends to associative-learning processes. It is well established that associative learning declines with age (Li, Naveh-Benjamin, & Lindenberger, 2005; Naveh-Benjamin, Craik, Guez, & Kreuger, 2005). Typical research paradigms involve forming associations between two random word pairs (e.g., *tree-bus; rose-sleep*) over multiple learning trials. Associative learning is an active process that draws on controlled cognitive abilities known to decline with age (Naveh-Benjamin et al., 2005). However, when prior knowledge can be accessed to support the learning of new associations, age differences are significantly reduced (Amer & Hasher, 2014; Castel, 2005).

This has also been investigated in the context of functional learning (Musielak, Giraudeau, Chasseigne, & Mullet, 2014). Functional learning is a form of associative memory that involves forming associations between item pairs (e.g., words) and a criterion item (e.g., a number). The critical adaptation from standard associative-memory paradigms is that the word pairs and criterion variable are organized along a common superordinate dimension (e.g., size) that can be abstracted and used to support associative learning. As an example, tree-elephant would be associated with a larger number, whereas flower-insect would be associated with a smaller number. In this form of associative learning, where access to prior knowledge can be used to support abstraction, age differences are practically eliminated, even in the oldest old (Musielak et al., 2014). Critically, functional learning has been argued to better reflect learning in the real world, where prior knowledge is frequently used to form new associative knowledge, in contrast to the learning of completely random associations common in most laboratory paradigms. As we saw above with the Moses and Truth Illusions, access to prior knowledge can also protect older adults from learning false associations in potentially manipulative situations such as misleading advertising (Brashier et al., 2017) or product pricing (Castel, 2005).

There is growing evidence that engaging one's semantic-knowledge stores in later life can lead to better cognitive performance even on tasks typically associated with cognitive-control abilities. For example, in a recent study, older adults showed equivalent learning to that of younger participants for a response set necessary for performance on a Stroop color-word test of inhibition. Critically, they were able to leverage this prior knowledge and transfer it across task conditions, resulting in equivalent performance between age groups on this classic test of cognitive-control functioning (Cohen-Shikora, Diede, & Bugg, 2018).

Recent evidence from our laboratory has shown that, despite declines in fluid cognitive abilities, older adults perform similarly to younger adults on a cognitive-control task assessing creative ability (divergent-thinking task; Adnan, Beaty, Silva, Spreng, & Turner, 2019). Note that

older people's performance on this task (but not younger people's) was associated with greater coupling of prefrontal and default-network brain regions (see below). This suggested to us that older adults may be relying on a relatively preserved store of semantic knowledge (associated with default-network function) to support performance on this task in the context of declining control resources.

Even incidental engagement of prior knowledge can support controlled cognition in older adulthood. In an interesting demonstration of this phenomenon, older and younger adults were asked to perform a standard Stroop task in which they were required to ignore irrelevant information (word meaning) and focus on the color in which the word was displayed. Despite explicit instructions to ignore the words, older, but not younger, adults showed encoding of the familiar (albeit distracting) word meanings, presumably supported by associations with prior semantic knowledge. Note that the incidental encoding of the distracting information led to better performance for older than younger adults on a subsequent general-knowledge test in which the previously distracting words were now task-relevant (Amer & Hasher, 2014).

# Multidomain abilities and real-world implications

The studies described above provide evidence that engaging prior knowledge, even incidentally, can both hinder and support performance across multiple domains, from memory and associative learning to cognitive-control tasks. The semanticization of cognition can also affect performance on more integrative, multidomain cognitive abilities such as problem solving. In a report involving older and younger adults, participants were asked to select the most advantageous choice from among an array of options to maximize financial gains and minimize losses in a decisionmaking task (Blanco et al., 2016). Unknown to the participants, the researchers imposed either a randomtrial architecture (i.e., previous knowledge could not predict future outcomes) or a structure such that previous response patterns could be used to make advantageous future choices. As predicted, older adults were slower in making decisions than the younger participants across both conditions. But they made significantly better decisions in the predictive condition, in which reliance on past knowledge was able to boost decision-making ability. However, they carried over this strategy, searching through previous knowledge stores, to the randomized trial blocks, resulting in poorer decisions for this condition. The authors concluded that a lifetime of reliance on prior knowledge, a typically adaptive strategy in real-world contexts, led older adults to access their knowledge stores, even when it was maladaptive to do so (Blanco et al., 2016).

The contribution of previous experience and knowledge to cognitive functioning has also been observed during real-world problem-solving tasks. Older adults demonstrate relatively greater reliance on prior knowledge during open-ended social problem-solving tasks. In contrast, younger adults rely to a greater extent on controlled, episodic recollection and simulation processes. These studies suggest that an age-related shift toward semanticized cognition may have a direct impact on how older adults approach and ultimately solve problems in their everyday lives (Sheldon, McAndrews, & Moscovitch, 2011; Vandermorris, Sheldon, Winocur, & Moscovitch, 2013).

The semanticization of cognition in older adulthood has also been associated with altered reasoning and decision-making capacity. Fuzzy-trace theory (Reyna & Brainerd, 1995) posits that, with increasing age, these complex fluid intellectual capacities increasingly rely on more crystallized, or gist-based, cognitive processing. Reliance on these "fuzzy," gist-based representations in older adulthood can be maladaptive, resulting, for example, in more false memories on the DRM paradigm as verbatim recollection declines (Brainerd & Reyna, 2015). However, gist-based processing, engaging lower-order and thus more flexible mnemonic representations, is also associated with more intuitive decision making as behavior becomes less constrained by fixed or verbatim depictions of past experience (Reyna, 2012). Consistent with the semanticization of cognition, gist-based memory is preserved relative to verbatim memory in older adulthood (Brainerd & Reyna, 2015; Brainerd, Reyna, & Howe, 2009; Reyna & Brainerd, 2011). Thus, decision making in real-world contexts, which requires more flexible, or gist-based, processing to adaptively map past experience to current contexts may be an area of relative strength in later life.

Consistent with this idea is the somewhat surprising performance of older adults, who performed as well as or better than younger adults on a series of economic decision-making tasks (Y. Li et al., 2013). The researchers assessed four aspects of decision making: temporal discounting, financial literacy, debt literacy, and loss aversion. Indeed, performance among older adults was strongly correlated with their crystallized (but not fluid) cognitive abilities, an association not observed for the younger cohort. These findings led the authors to propose the *complementary cognitive capabilities hypoth*esis, which states that access to an expanding and stable repertoire of stored knowledge can support cognitive performance and may even eliminate age-related cognitive decline when prior knowledge is accessible, accurate, and adaptive for the current task context.

This shifting architecture of cognition across the life span, from cognitive-control abilities to greater reliance on prior knowledge, has also been used to explain the economic decision-making "sweet spot" in middle age (Agarwal et al., 2009; Samanez-Larkin, 2013). In this model of real-world financial decision making, late 40and 50-year-olds, falling roughly at the intersection of declining control and increasing semantics (Park et al., 2001; see also Fig. 1), retain the capacity to flexibly adjust to shifting economic contingencies, while also drawing on an accumulating repertoire of past experience, to support more adaptive financial borrowing strategies. This idea was directly tested in a study that compared the impact of fluid and crystallized cognitive abilities on financial credit scores (Y. Li et al., 2015). Domainspecific crystallized knowledge (i.e., financial literacy) was able to offset the loss of fluid cognitive abilities and predicted higher credit scores in later life.

Moving beyond the financial domain, further support for the influence of prior knowledge on decision making has been demonstrated as greater regard for intertemporal choice (Lockenhoff, 2011). This simply refers to the amount of value discounting that occurs when deciding between an immediate versus a larger delayed reward. Younger adults show a steep discounting rate, exhibiting a strong preference for the immediate reward. In contrast, older adults show a much shallower rate of discounting, resulting in a similar valuation of immediate and delayed rewards. This suggests that older adults can bring their lifetime of experience and knowledge about the benefits of postponed gratification to bear on their economic choices. Research from the emerging field of decision neuroscience is beginning to elucidate the neural mechanisms underlying the influence of prior knowledge on decision-making processes in older adults (Samanez-Larkin & Knutson, 2014, 2015).

In this section we reviewed evidence that the semanticization of cognition has a profound impact (both positive and negative) on cognitive and real-world functioning in later life. Next we explore the determinants of this shift in cognitive architecture from younger to older adulthood. We examine whether these changes reflect an enduring shift in cognitive mode and whether increased reliance in later life on prior knowledge is an artifact of declining control abilities or a more deliberate shift in motivation toward favoring familiarity over novelty.

# Increasing Semantics and Declining Control: A Shift in Cognitive Mode in Older Adulthood?

In the previous section, we described how the shifting architecture of cognition can both benefit and harm performance across multiple task contexts. Here we turn to address more directly the question of why such a change occurs. What are the central determinants of greater reliance on prior knowledge to guide thought and action in later life? We explore whether the accrual of semantic knowledge across the life span is alone sufficient to precipitate this shift or whether a decline in cognitive control is necessary to promote the semanticization of cognition. We then address whether the semanticization of cognition is an in situ, or contextdependent, response or perhaps reflects a more durable shift in cognitive mode. Finally, we discuss socioemotional selectivity theory as an alternate account, suggesting that semanticized cognition in later life may not merely be a reaction to shifting cognitive resources but rather an active strategy to promote affective well-being in the context of a foreshortened time horizon.

### Increasing semantic knowledge

As should be clear from the previous section, the shifting architecture of cognition in later life conveys both benefits and costs with respect to task performance across multiple cognitive domains. Investigating how and when prior knowledge is brought on-line to influence cognitive functioning may provide important insights into the shifting nature of cognitive and realworld functional abilities in later life. Crystallized, or semantic, knowledge is accessed more readily, drawing on fewer attentional resources than new learning, which involves cognitive control to form novel associations between previously discrete percepts or between percepts and prior knowledge (Craik & Jennings, 1992). Through repeated exposure, reencoding, and consolidation, activation of prior-knowledge structures on the basis of stimulus familiarity becomes less attention demanding and increasingly automatic (Naveh-Benjamin et al., 2005). As a result, the accumulation of prior knowledge in the form of facts, habits, routines, schema, and even stereotypes or prejudices exert greater control as familiarity increasingly drives cognition in later life.

Increases in semantic knowledge across the life span suggests that the intrusion of prior knowledge into ongoing thought and action will increase simply as a function of its availability. A larger store of more frequently encoded semantic knowledge increases the probability that it will be activated by specific percepts or thoughts occurring within a given task context. As older adults possess a larger store of semantic knowledge (Hoffman, 2018; Verhaeghen, 2003), it follows that knowledge representations will be activated across an ever larger range of perceptual and mental contexts. But is knowledge accumulation sufficient to explain the breadth of influence semantics exert on cognitive functioning in later life? Work suggests that declining control processes also play a critical role.

### Declining cognitive control

Analogous to the separation of intellectual ability into fluid and crystallized capacities, the domain of semantic cognition has been similarly divided into semantic and control processes that facilitate access to a growing store of prior knowledge in later life (Hoffman, 2018; Jefferies, 2013; Ralph et al., 2017). In this model, "semantic control" is differentiated from domain-general cognitive-control processes. Semantic control increases with age and enables older adults to strategically retrieve and monitor access to weaker semantic associations embedded within an expanding store of semantic knowledge. Put another way, semantic-control processes, honed over time to strategically search this expanding database, can better leverage semantics to influence cognition (Hoffman, 2018). This idea is consistent with evidence, reviewed earlier, that older adults can draw more readily from prior knowledge to make better, albeit slower, decisions than younger adults (Blanco et al., 2016).

However, gains in semantic control occur in the context of declining domain-general control processes, such as inhibitory processing (e.g., Hasher & Zacks, 1988). Reduced cognitive control can lead to poor filtering of irrelevant semantic associations and the leakage of prior knowledge into ongoing cognitive operations. Evidence for this "leaky-semantics" account is found in the false-memory studies reviewed earlier (e.g., Koutstaal et al., 2003; Schacter, Koutstaal, Johnson, Gross, & Angell, 1997; Schacter, Koutstaal, & Norman, 1997). In an ostensibly more adaptive context, the intrusion of semantics in the form of incidental encoding of familiar (but distracting) information, has been shown to improve performance on a subsequent task for which previously distracting information had become relevant to current task goals (Amer & Hasher, 2014). Age-related declines in domain-general cognitive control may, perhaps counterintuitively, also lead to poorer engagement of prior knowledge. In an investigation of semantic fluency, older adults displayed more frequent switching between retrieval cues, leading to poorer performance on a semantic-fluency task (Hills, Mata, Wilke, & Samanez-Larkin, 2013). The authors interpreted this finding as evidence that declining inhibitory control in older adulthood resulted in an inability to suppress distracting retrieval cues, leading to increased and less efficient switching during semantic retrieval.

Together, these findings identify declining domaingeneral control processes as an important determinant of the shift toward increasingly semanticized cognition in older adulthood. Specifically, there is mounting evidence that declining inhibitory capacity in older adulthood enables greater intrusion of prior knowledge into ongoing cognitive operations. This suggests that the semanticization of cognition is not solely determined by an accumulation of prior knowledge. Reduced cognitive-control abilities also contribute to the growing influence of semantics on cognitive processing in later life

### A cognitive-mode bypothesis

This two-factor account of the semanticization of cognition in older adulthood, implicating a larger repertoire of semantics and declining control processes, parallels an emerging theory of goal-directed action that hypothesizes that human behavior is shaped by a constant tension between exploratory versus exploitative modes of functioning (Hills et al., 2015). Exploratory behaviors include a preference to seek novel associations, an orientation to learning, and a focus on the external environment over prior knowledge. Exploitative behaviors involve a preference for relying on previous knowledge and expectations and a general aversion to novelty (Baror & Bar, 2016; Hills et al., 2015; Schwartenbeck, Fitzgerald, Dolan, & Friston, 2013).

Within the domain of human cognition, availability of cognitive-control resources is an important factor in the shift from an exploratory to a more exploitative cognitive mode (Baror & Bar, 2016). In this study, younger adults were asked to generate free associations under conditions of high and low attentional load. Participants generated a greater number of novel, or exploratory, associations under low-load conditions compared with the high-load condition. As attentional resources were increasingly taxed, associations became more uniform, or exploitative, in nature. Further, subsequent priming of close associations was observed for high-load conditions and for more remote associations under low attentional load. These results provide strong evidence that the availability of attentional resources determined whether participants approached the task from an exploratory or an exploitative cognitive mode (Baror & Bar, 2016). Lower cognitive-control resources resulted in a shift toward the exploitation of prior knowledge and away from novelty seeking or the generation of new associations.

Although these results have yet to be replicated in an older adult sample, the prediction for cognitive aging is clear. Declines in cognitive control, a hallmark of cognitive aging, should result in a shift from exploration to exploitation as a cognitive mode in older adulthood (see Fig. 2). There is early, albeit indirect, support for this idea. Under conditions of high attention load, older adults are more likely to produce false memories on the DRM paradigm (Koutstaal, Schacter, & Bernner, 2001), suggesting greater influence of prior knowledge

on mnemonic performance. Likewise, the relative advantage for older adults on functional versus associative memory tasks has been attributed to the coupling of new and prior knowledge in working memory (Musielak et al., 2014). There is also evidence that agerelated declines in episodic recall, associated with reduced control processes, results in an age-related shift toward exploiting prior knowledge as older adults increasingly draw on semanticized recall to support memory functioning (Levine et al., 2002; Spaniol & Bayen, 2002). A shift in cognitive mode would also be consistent with our findings on creativity in aging discussed earlier. Older adults showed greater coupling of default-executive control brain regions during both task (Adnan, Beaty, Silva, Spreng, & Turner 2019) and rest (Adnan, Beaty, Lam, Spreng, & Turner 2019) and this coupling was associated with better divergent thinking task performance across our sample of older adults. We interpreted these findings as older adults showing greater reliance on prior knowledge, or an exploitative cognitive mode, during creative cognition.

## Semanticization of cognition and socioemotional selectivity

Evidence reviewed thus far suggests that a shift toward the semanticization of cognition, or an exploitative processing mode, occurs in response to a growing pool of semantic knowledge and reduced cognitive control in older adulthood. However, this shift may not merely be a reaction to the altered balance of knowledge and control but rather an active, intentional process precipitated by a shifting motivational hierarchy in later life. Socioemotional selectivity theory (SST; for a review, see Carstensen et al., 2003) suggests that the perception of a foreshortened time horizon shifts goal structures in older adulthood away from more expansive pursuits of new knowledge (i.e., an exploratory mode) and toward emotionally meaningful goals that provide more immediate personal satisfaction (i.e., an exploitative mode). According to SST, the shift from exploration to exploitation in later life reflects a parallel shift in motivational drives resulting from an increasing awareness of the finiteness of time. Perceived shorter time horizons motivate greater focus on more immediate and affectively salient goals.

Among the earliest evidence for SST was the discovery that the balance of one's social networks shifted toward increasingly close or intimate relationships in later life, with a relative decrease in the engagement of new or more casual acquaintances (Carstensen, 1992). These data suggest that the shift toward exploiting prior experiences extends to the domain of social functioning, manifest in this context as a greater focus on more

intimate social relationships. Critically, this socioemotional explanation for an age-related shift toward an exploitative cognitive mode is consistent with our DECHA model (discussed in detail below). A core node of the default network, the medial prefrontal cortex, and its connectivity to lateral prefrontal brain regions implicated in cognitive control, has been associated with many aspects of social cognition (Amodio & Frith, 2006; Forbes & Grafman, 2010). Critically, our work has also shown that this medial-to-lateral prefrontal-cortex connectivity pattern is greater in older versus younger adults during both task (Turner & Spreng, 2015) and rest (Spreng et al., 2018), consistent with DECHA.

In this section, we demonstrated that the ongoing accrual of semantic knowledge and experience across the adult life span, in the context of declining control resources, leads to the increasing semanticization of cognition and a shift in cognitive mode, from exploration to exploitation in later life. However, the neural basis of the shifting architecture of cognition has yet to be fully explored. In the following section we shift the focus of our review to the aging brain, highlighting the patterns of functional brain changes that we believe closely parallel the shift in cognitive architecture described in these opening sections. In the final section of the review we integrate these brain changes into a single model, DECHA, a candidate neural mechanism bridging life-span changes in cognitive and brain aging.

# The Shifting Architecture of Brain Structure and Function in Aging

In the prior section, we discussed how the accrual of knowledge and experience across the adult life span, along with declining control resources, leads to the increasing semanticization of cognition with advancing age. We now turn to an examination of the shifting architecture of the brain that may underlie this co-occurring shift in cognitive architecture. We review structural and functional brain changes broadly before focusing on changes specific to lateral prefrontal-cortex and default-network modulation with advancing age.

### Structural brain changes

The human brain undergoes significant structural and functional changes across the adult life span (Spreng & Turner, 2019). Global changes in gray matter, white matter, and ventricular volumes are a hallmark of normal brain aging. Lateral prefrontal cortices, implicated in cognitive control, appear particularly vulnerable to age-related structural change (Raz et al., 2005). Changes to the brain's white matter, axonal projections supporting communication among spatially distributed regions,

or networks, are also a prominent feature of brain aging and are a strong predictor of age-related cognitive decline (Piguet et al., 2009; Snowdon, 1997; Yarchoan et al., 2012). Among large-scale brain networks, structural changes to the default network have been commonly reported (Andrews-Hanna et al., 2014). Structural covariance studies have demonstrated network-level changes in typical aging, and the reduced integrity of this network is considered a neural network biomarker of the transition from typical to pathological aging (Spreng & Turner, 2013).

### Functional brain changes

The relationship between structural and functional brain changes remains an area of active debate and inquiry. Both the STAC (Park & Reuter-Lorenz, 2009) and CRUNCH (Reuter-Lorenz & Cappell, 2008) theories of neurocognitive aging suggest that structural brain changes may lead to the promotion of new, compensatory functional connections. Among the most commonly reported brain changes are greater recruitment and poor modulation of neural activity, primarily in prefrontal brain regions, in response to changing task demands (Cabeza, 2002; Grady, 2012). More globally, aging is associated with a pattern of functional brain change commonly referred to as neural dedifferentiation, or an inability to recruit specialized neural circuits associated with discreet processing operations (Park et al., 2001). Whether this dedifferentiation reflects a compensatory or maladaptive process remains a point of some debate (Grady, 2012; Spreng & Turner, 2019; Turner & D'Esposito, 2010).

As we noted at the outset, functional brain changes are increasingly investigated through the lens of brain networks, or functionally connected assemblies of spatially distributed brain regions. Paralleling local changes, brain networks also become increasingly dedifferentiated in older adulthood. Network-level changes are marked by reduced within-network and increased between-network connections (e.g., Chan, Park, Savalia, Petersen, & Wig, 2014; Geerligs, Renken, Saliasi, Maurits, & Lorist, 2015). These network interactions are also poorly modulated by task demands (Damoiseaux, 2017). To highlight the parallels between cognitive and functional brain aging, here we focus in more detail on two of the most commonly reported patterns of agerelated functional brain change: increased recruitment of lateral prefrontal brain regions and reduced suppression of the default network.

**Lateral prefrontal cortex activation.** One of the earliest and most persistent patterns of age-related functional brain change observed during goal-directed tasks is enhanced bilateral recruitment of lateral prefrontal cortices (as in

HAROLD; Cabeza, 2002). Lateral prefrontal cortices, and connections with posterior and subcortical brain regions, are critical for implementing cognitive-control processes (Duncan, 2010; Shallice & Burgess, 1996; Stokes et al., 2013; Stuss & Levine, 2002). As cognitive-control decline is a central feature of cognitive aging (Verhaeghen & Cerella, 2002), increased engagement of prefrontal cortices may reflect greater control demands in older versus younger adults.

This idea, most directly captured in the CRUNCH (Reuter-Lorenz & Cappell, 2008), posits that, with age, structural and functional brain changes result in noisier, lower-fidelity, and poorly gated inputs for cognitiveprocessing operations (Gazzaley, Cooney, Rissman, & D'Esposito, 2005; S. C. Li & Rieckmann, 2014; Schmitz, Dixon, Anderson, & De Rosa, 2014). These degraded inputs necessitate greater cognitive control, mediated by lateral prefrontal brain regions, to extract goalrelevant signals from this noisier cognitive landscape (Park & Reuter-Lorenz, 2009). However, enhanced prefrontal recruitment may also be maladaptive, reflecting a loss of functional specialization (or dedifferentiation) secondary to structural brain changes (Park et al., 2001). In this interpretation, aging is associated with the degradation of specialized neural circuits, resulting in more diffuse and less efficient neural processing. While the cognitive implications continue to be debated, both compensation and dedifferentiation accounts converge in their prediction of age-related increases in lateral prefrontal brain activity during task performance.

**Default-network suppression.** With the advent of whole-brain, functional neuroimaging methods more than 2 decades ago, changes in large-scale functional brain networks are rapidly becoming a hallmark of functional brain aging (Damoiseaux, 2017; Grady, 2012). As we noted earlier, the default network is an assembly of functionally connected brain regions, including the posterior cingulate cortex, medial prefrontal cortex, inferior parietal lobule, and medial and lateral temporal lobes (Buckner, 2004). This network was first identified for its reliable suppression across a range of externally directed tasks in young (Raichle et al., 2001) and older (Buckner et al., 2008) adults. Although its functions continue to be debated (Spreng, 2012), the default network is activated during internally directed cognitive processes, including access to stored knowledge representations and experiences (Andrews-Hanna et al., 2014). Critically, this network has been implicated in associative and elaborative processing (Bar, Aminoff, Mason, & Fenske, 2007), linking perceptual information with internal representational knowledge structures to support cognitive processing (Spreng et al., 2014).

The default network is vulnerable to age-related changes. Activity within this network, typically suppressed

during externally directed cognitive tasks in young adults, is less deactivated in older adults (Lustig & Buckner, 2004). Functional connectivity within the default network also declines with age (Andrews-Hanna et al., 2007). Work suggests that reductions in connectivity within the default network predicts greater connectivity with frontoparietal control regions (Grady, Sarraf, Saverino, & Campbell, 2016). Critically, this pattern of between-network connectivity appears to be poorly modulated by task context, with greater cross talk observed at lower levels of task demand and across task contexts for older versus younger adults (Spreng & Schacter, 2012; Turner & Spreng, 2015). Poor modulation of network interactivity on the basis of task context represents a fundamental alteration in the network architecture of the brain. As we argue in the next section, greater, and less flexible, coupling of default and prefrontal brain regions (i.e., DECHA) may provide a key neural conduit promoting the semanticization of cognition in later life.

Here we have focused on alterations in cortical structure and function to provide the necessary foundation for the introduction of our DECHA proposal in the following section. However, brain aging encompasses a much wider array of changes, including alterations in the structure and function of subcortical structures and neurotransmitter systems. Although a more comprehensive review of these changes is beyond the scope of the review, one particular theoretical account of dopamine signaling, novelty-related motivation of anticipation and exploration by dopamine (NOMAD; Duzel, Bunzeck, Guitart-Masip, & Duzel, 2010), is of particular relevance to our model of semanticized cognition, or the emergence of an exploitative cognitive mode in later life. In the NOMAD account, reduced dopamine signaling within striatal-hippocampal circuitry is associated with age-related reductions in exploratory or novelty-seeking behaviors. This suggests that as striatal dopamine functioning declines with age, reduced dopaminergic tone within striatal-hippocampal circuits may impede cognitive processes associated with exploration, including episodic simulation, prospection, and planning for future events. Thus, a prominent feature of brain aging, reductions in the structure and function of the hippocampus (Buckner, 2004), mediated in part by deficient dopaminergic signaling, may be a neural mechanism underlying the shift toward a more exploitative cognitive mode in older versus younger adults. The hippocampus and the medial temporal-lobe formation make up one subsystem of the default network (Andrews-Hanna, Reidler, Sepulcre, Poulin, & Buckner, 2010). Although speculative, the loss of dopaminergic signaling to the hippocampus may alter the functional integrity of the default network more broadly, thereby precipitating greater coupling with frontal control networks, and promote a more exploitative cognitive mode. Indeed, this pattern of default-frontal coupling in response to declines in the functional integrity of the default network has been observed (Grady et al., 2016).

In the final section of the review, we further explore the neural mechanisms associated with the semanticization of cognition and a more exploitative mode of cognitive processing. Specifically, we argue that DECHA provides a critical mechanism, linking the broad patterns of cognitive and brain aging discussed thus far in the review.

# DECHA: A Unified Account of Cognitive and Brain Aging

Although age-related changes to lateral prefrontal and default-network brain regions are both hallmarks of neurocognitive aging (Grady, 2012), there has been scant consideration of whether or how these patterns intersect until recently. We demonstrated that lateral prefrontal engagement and reduced default-network suppression co-occur and are functionally coupled in older adulthood, leading us to propose DECHA (Turner & Spreng, 2015). We hypothesize that these functional brain changes parallel the shift in cognitive mode, from declining control processes, reflected in poorly modulated lateral prefrontal cortex activation, to greater reliance on prior knowledge, reflected in reduced default-network suppression during goal-directed tasks. We suggest that functional coupling of these two activity patterns provides a neural conduit promoting the semanticization of cognition (see Fig. 2).

### DECHA as dedifferentiation?

Network dedifferentiation, resulting in greater cross talk between brain networks, is a common feature of functional brain aging and has been generally associated with age-related cognitive decline (Chan et al., 2014; Geerligs et al., 2015). In this context, DECHA, as a specific exemplar of network dedifferentiation, should be associated with poorer cognitive performance in older versus younger adults. Indeed, we observed this in our study of the Tower of London task (Turner & Spreng, 2015), a measure of externally directed planning that does not require access to stored representations (Shallice, 1982). Moreover, greater default-executive coupling has been associated with lower fluid intellectual functioning during both task (Rieck et al., 2017) and rest (Spreng et al., 2018). However, a study examining age-related changes in functional connectivity among three networks specifically implicated in goal-directed cognition, including default, attention, and cognitive-control networks, suggests a more complex picture (Grady et al., 2016).

Functional connectivity within and between networks was examined during the performance of three tasks (associative memory, selective working memory, and trait judgment). Between-network connectivity of the cognitive-control network (including core nodes in the lateral prefrontal cortex) increased with age, and this pattern was associated with better performance on the associative-memory and trait-judgment tasks in older, but not younger, adults. Interestingly, associations were not observed for the selective working memory task, which, unlike the associative-memory and trait-judgment tasks, did not rely on mnemonic processing or activation of stored knowledge representations. Thus, consistent with DECHA, increased connectivity between frontal and default-network brain regions can lead to better cognitive performance. However, as we saw earlier, this benefit depends on the congruence between task goals and prior knowledge.

#### **Context matters**

An important early test of this principle was reported by Spreng and Schacter (2012). In this study, younger and older adults performed the Tower of London planning task and an analogous autobiographical future planning task. The two tasks were closely matched for sensorimotor demands, and both involved a planningdomain manipulation. Unlike the Tower of London task, the autobiographical planning task required participants to access personally meaningful information to generate future plans. Both older and younger adults showed coupling of default and frontoparietal control networks during the autobiographical planning task, consistent with the idea that default-executive coupling can support cognition when prior knowledge is congruent with task goals. During the Tower of London task, the functional connectivity patterns for younger and older participants diverged. Whereas younger adults showed greater coupling between frontoparietal-control and visual-attention regions, consistent with performing an externally directed visual task, older adults failed to decouple lateral prefrontal and default-network brain regions. In short, they failed to modulate functional connectivity between default and executive-control regions on the basis of task context. This study provided the first direct evidence of task-based affiliation between default and executive-control regions during a goal-directed task for which access to prior knowledge was relevant (i.e., autobiographical planning). Further, for young adults, network affiliations shifted between the two planning tasks on the basis of the relevance of prior knowledge, whereas older adults failed to flexibly shift network connectivity on the basis of task context. This suggests that default-executive coupling may be less flexible in older adulthood. But how do these network-level findings relate to the agerelated shift in cognitive modes we discussed earlier? As we saw, when planning was impersonal during the Tower of London task, young adults decoupled default and frontal brain regions; older adults maintained default-executive coupling and performed more poorly on the task. Goal relevance of prior knowledge and age are therefore key determinants of whether DECHA reflects an adaptive or maladaptive process.

When tasks require processing of stimuli devoid of personal meaning and unconnected to prior knowledge, DECHA predicts that positive coupling between default and executive-control regions will lead to poorer performance in older adults. There is growing evidence for this idea. In a life-span study of brain activity during performance on a spatial-reasoning task, greater activation of lateral prefrontal cortex and greater deactivation of the default network in response to increasing task demand were correlated. Further, this association between positive and negative modulation in activity predicted poorer task performance as well as lower fluid-reasoning ability outside of the scanner across the life span (Rieck et al., 2017). These findings are consistent with several studies showing that reduced suppression of the default network and increased coupling with lateral prefrontal cortices is associated with poorer performance on externally directed cognitive tasks (Damoiseaux, 2017; Grady, 2012; Keller et al., 2015; Persson, Lustig, Nelson, & Reuter-Lorenz, 2007; Spreng, Stevens, Viviano, & Schacter, 2016; Turner & Spreng, 2015).

Few studies to date have investigated how defaultexecutive coupling influences cognitive performance when access to prior knowledge is consistent with task goals. It has long been assumed that default-network activity must be suppressed to facilitate performance on all cognitive-control tasks. However, studies have challenged this idea. Using a social-working memory paradigm, Meyer, Spunt, Berkman, Taylor, and Lieberman (2012) asked participants to hold in mind personality traits of known others. They observed robust activation of core default-network regions typically implicated in social-cognitive tasks. Critically, this default-network activity co-occurred with the activation of lateral prefrontal cortices that are implicated in working memory processes. Although the authors did not report functionalconnectivity analyses, this study provided early evidence that the default network is not uniformly suppressed during cognitive-control tasks and that default and lateral prefrontal brain regions can be simultaneously engaged when cognitive-control operations require access to stored knowledge representations (e.g., Baird, Smallwood, Gorgolewski, & Margulies, 2013; Konishi, McLaren,

Engen, & Smallwood, 2015; Spreng et al., 2014; Spreng, Gerlach, Turner, & Schacter, 2015; Spreng, Stevens, Chamberlain, Gilmore, & Schacter, 2010).

## DECHA as an enduring shift in functional brain aging

So far, we have shown that the behavioral impact of default-executive coupling, as a putative mechanism underlying the semanticization of cognition in later life, depends on the congruence between prior knowledge and task goals. The studies reviewed thus far relied on different tasks to demonstrate this point. However, a critical prediction of DECHA is that greater default-executive coupling should be associated with a shift in cognitive processing toward greater reliance on semantics in older adulthood. Demonstrating such a relative shift in the influence of prior knowledge across the life span requires measuring the relative contributions of cognitive control and semantics within a single task.

To test this prediction we investigated autobiographical memory performance and functional network architecture in younger and older adults (Spreng et al., 2018). We used the Autobiographical Interview (Levine et al., 2002), which enabled us to generate separate estimates of episodic and semantic recollection. Deriving these two measures from a single task allowed us to contrast the level of controlled retrieval, necessary for episodic recall, with the level of semanticized knowledge generated during remembrances of personal past events. Default-executive coupling was assessed with resting functional MRI. We reasoned that if the shifting architecture of cognition occurs across the adult life span, then this shift should be most robustly reflected in these intrinsic network-connectivity patterns, which are also shaped throughout the life course (Stevens & Spreng et al., 2014). We first observed that default-executive coupling at rest was negatively associated with a measure of fluid intelligence. This was consistent with our argument that default-executive coupling is not adaptive when prior knowledge is incongruent with task goals. The second and more critical finding with respect to DECHA was that greater default-executive coupling was associated with more semanticized autobiographical retrieval for older, but not younger, adults (even when controlling for fluid cognition). These results provide support for the idea that greater default-executive coupling is associated with the semanticization of cognition in later life (Spreng et al., 2018). Further work is necessary to test this hypothesis in additional cognitive domains in which age-related shifts toward semanticized cognition have been reported. In one notable study, older adults showed greater default network activity and enhanced functional connectivity between default and controlnetwork regions during retrieval of realistic versus unrealistic grocery-item prices using a subsequent memory paradigm. The authors interpreted these functional brain differences as evidence for greater reliance on stored memory representations for meaningful associations in older adults in contrast to greater reliance on controlled encoding and retrieval processes in young (Amer, Giovanello, Nichol, Hasher, and Grady, 2019).

# Determinants of default-executive coupling

To further explicate DECHA as a mechanism for predicting age-related cognitive changes, we need to better understand the determinants of default-executive coupling and how variations in this pattern of network interactivity are associated with the semanticization of cognition. We have identified two factors: congruence with prior knowledge and the availability of control resources. As we reviewed earlier, familiarity drives the semanticization of cognition. At the level of the brain, default-network brain regions are activated simply by perceiving familiar stimuli (Bar et al., 2007; Leveroni et al., 2000; Spreng et al., 2014). Engagement of the default network by known stimuli, in the absence of explicit task demands, is consistent with a knowledgebased account of the semanticization of cognition in later life. Age-related increases in the accumulation and fluency of stored representations (Craik & Jennings, 1992) would result in greater default-network activation and influence of prior knowledge on cognitive functioning in older versus younger adults. As we have seen, this can be both adaptive, as in the Moses and Truth Illusion experiments, or maladaptive, as in the falsememory studies. However, activation of prior knowledge is a necessary but not sufficient determinant of default-executive coupling. Goal-directed tasks additionally require activated knowledge representations to enter the focus of attention to influence future thought and action. As access to the focus of attention is modulated by cognitive-control processes, the availability of control resources is a second determinant of semanticized cognition in later life.

The interdependence of prior knowledge and control processes during ongoing cognitive operations was convincingly demonstrated in the investigation of exploratory and exploitative modes of cognitive processing described earlier (Baror & Bar, 2016). In that study, cognitive control was supplanted by access to prior knowledge to support task performance. Likewise, as control resources decline with age, prior-knowledge

representations would be expected to move to the fore and exert greater influence over cognitive operations. We argue that greater, and less flexible, default-executive coupling in older adults serves as a putative neural mechanism associated with greater semanticization of cognition (Fig. 2).

In a review of brain activity during semanticprocessing tasks, Binder, Desai, Graves, and Conant (2009) identified a robust pattern of default and lateral prefrontal activation in young adults. A recent review contrasting brain activity between younger and older adults identified age differences in lateral prefrontal brain cortex but not in the default network (Hoffman & Morcom, 2018). As we have seen throughout the review, accessing prior knowledge conveys larger performance benefits for tasks that more closely mirror real-world functions, when past patterns of thought and action can be accessed to support current or planned behavior. However, there are two important differences between real-world and laboratory-based tasks. As we discussed earlier, laboratory stimuli are typically devoid of personal relevance, rendering knowledge of one's past behavior at best irrelevant and at worst distracting. In addition, laboratory-based tasks are highly constrained. Task rules and the current stimulus array, by necessity, guide behavior. Here again, knowledge of past behavior is unnecessary or potentially maladaptive.

Thus, the failure to observe robust, age-related changes in the default network during standard laboratory-based tasks of semantic cognition is perhaps unsurprising. As we have shown, engaging prior knowledge is more beneficial during less constrained, more real-world task contexts such as economic decision making, problem solving, or future planning. Critically, older adults display this tendency more than their younger counterparts, whether or not they are instructed to do so (Blanco et al., 2016; Musielak et al., 2014). Under these task conditions, mimicking unconstrained real-world contexts, increased activation, and coupling of the default network with lateral prefrontal regions may not reflect maladaptive dedifferentiation but rather a change in cognitive mode secondary to declining control resources with age (c.f. Baror & Bar, 2016). Constraints imposed by the task, precluding engagement or leveraging of this exploitative cognitive mode, may be another important determinant of default-executive coupling in later life. Unconstrained or everyday tasks leave greater scope to search and engage prior knowledge. Under DECHA, this would in turn lead to greater default-network activity and increased coupling with lateral prefrontal brain regions to grant these representations privileged, or at least priority, access to the focus of attention.

We have now identified three putative determinants promoting the semanticization of cognition in older adulthood: congruence between items in the focus of attention and prior knowledge (i.e., familiarity), availability of cognitive-control resources, and the nature of the task, either promoting or precluding an exploitative cognitive mode. Each of these determinants implicate either default or lateral prefrontal brain regions, or their interaction. Default-network activity is modulated by stimulus familiarity, whereas lateral prefrontal brain regions mediate cognitive-control processes and are most reliably engaged during unconstrained tasks. Although our work exploring DECHA as a neuralnetwork mechanism of cognitive aging is ongoing, we predict that a change in any of these factors (increased stimulus familiarity, decreased cognitive control, unconstrained task environment) would result in greater default-executive coupling and the semanticization of cognition in older adulthood.

# DECHA: a novel account of neurocognitive aging

It is important to reemphasize here that default-executive coupling, as reflected in DECHA, does not predict task performance per se. Rather, it is the context-dependent engagement and flexible coupling of lateral prefrontal and default brain regions that may best predict adaptive, goal-directed behavior in older adulthood. We suggest that it is the context dependency of DECHA that distinguishes it from other leading neurocognitive aging theories such as HAROLD (Cabeza, 2002), PASA (Davis, Dennis, Daselaar, Fleck, & Cabeza, 2008), or the CRUNCH (Reuter-Lorenz & Cappell, 2008) and STAC (Park & Reurter-Lorenz, 2009). By incorporating both dimensions of the shifting architecture of cognition within a single framework, DECHA can generate predictions of both cognitive gains and losses in later life. Further, we argue that considering both cognitivecontrol demands and the relevance of prior knowledge is particularly important for predicting older adult capacity in real-world settings, in which access to prior knowledge and experience may convey greater benefit than in typical laboratory-based contexts.

Finally, as reviewed above, results from our lab and others suggest that these age-related network changes are measurable at rest, in the absence of explicit task demands (Ng, Lo, Lim, Chee, and Zhou, 2016; Spreng et al., 2018). We have previously suggested that functional connectivity patterns such as default-executive coupling, entrained over the life span, may alter the intrinsic network architecture of the brain (Stevens & Spreng, 2014). These intrinsic network changes, shaped over years or decades, provide important insights into similarly protracted but durable shifts in cognitive abilities. We suggest that evidence for greater coupling

between the default-network and executive-control regions is fully consistent with an ontogenetic shift toward increasingly semanticized cognition and an exploitative cognitive mode in later life. However, within-subject studies contrasting task and resting-state network organization across age cohorts will be necessary to address whether poor task-driven modulation of default-executive coupling is an emergent property of a more enduring age-related shift in the intrinsic network architecture of the brain.

In this final section of the review we argue that DECHA provides not simply a descriptive but a mechanistic account of neurocognitive aging. By incorporating changes in brain regions associated with both cognitive-control processes (lateral prefrontal cortex) and semantic cognition (default network), DECHA may help to predict the dual trajectories of cognitive aging and further our understanding of how these cognitive and neural changes intersect to promote the shift from cognitive exploration to cognitive exploitation in later life.

#### Conclusion

Psychological and neuroscience studies of cognitive aging are disproving the fallacy that cognitive aging equals cognitive decline. The true story of cognitive aging is one of both gains as well as losses. With age comes an expanding repertoire of knowledge and life experience that can help navigate the routines—and the surprises—of daily life. Of course, repeating the past does not necessarily portend an optimal future. When and how past knowledge is accessed to guide goal-directed behavior can influence how successfully older adults can reason, solve problems, and plan in their everyday lives. Indeed, this idea of engaging prior knowledge to guide real-world behavior closely resembles definitions of wisdom, in which adaptive thought and action require access to tacit knowledge about oneself, others, and situational contexts (Sternberg, 1998). Better access to a richer store of tacit knowledge, or the fundamental pragmatics of life (Baltes & Staudinger, 2000), is considered to be the basis of practical intelligence and a precondition for wise thinking. Accessing these stores of prior knowledge, including representations of oneself, the thoughts and intentions of others, and how they interact in situ, all engage default-network brain regions (Andrews-Hanna et al., 2014). Although speculative, this raises the intriguing possibility that increased default-executive coupling may serve as a neural conduit allowing prior knowledge to enter the focus of attention and facilitate wise thinking in later life.

We have drawn on findings from cognitive psychology and cognitive neuroscience to develop a comprehensive account of neurocognitive aging. Integrating parallel shifts in the architectures of cognition and brain function, DECHA encompasses the broad spectrum of aging cognition, generating predictions for both gains and losses, and their interactions, across the life span. However, in contrast to the rich literature identifying the shift in cognitive architecture from young, to middle, to older age (e.g., Park et al., 2001; Fig. 1), few studies have directly examined life-span changes in functional brain-network interactions (but see Rieck et al., 2017; Chan et al., 2014). Future research is necessary to map interactions among functional brain networks and how they change across the full adult life course. A significant challenge going forward will be to identify a set of determining conditions for defaultexecutive coupling, and the semanticization of cognition, from younger to older adulthood. This will require novel paradigms to experimentally manipulate the engagement of prior knowledge to determine a putative "set point," marking the shift from exploratory to exploitative processing and investigating how this baseline changes across the life span. This would provide the necessary behavioral markers to further test DECHA and explore its utility as a neural mechanism to predict individual differences in the trajectory of cognitive and real-world functional capacities across the adult life span.

#### **Action Editor**

Timothy McNamara served as action editor for this article and June Gruber served as interim editor-in-chief.

#### ORCID iD

R. Nathan Spreng https://orcid.org/0000-0003-1530-8916

#### **Declaration of Conflicting Interests**

The author(s) declared that there were no conflicts of interest with respect to the authorship or the publication of this article.

#### References

Adnan, A., Beaty, R., Lam, J., Spreng, R. N., & Turner, G. R. (2019). Intrinsic default-executive coupling of the creative aging brain. *Social Cognitive and Affective Neuroscience*, *14*, 291–303.

Adnan, A., Beaty, R., Silva, P. J., Spreng, R. N., & Turner, G. R. (2019). Creative aging: Functional brain networks associated with divergent thinking in older and younger adults. *Neurobiology of Aging*, 75, 150–158.

Agarwal, S., Driscoll, J. C., Gabaix, X., & Laibson, D. (2009). *Age of reason: Financial decisions over the life cycle and implications for regulation*. Washington, DC: Brookings Institute.

- Amer, T., Giovanello, K. S., Grady, C. L., & Hasher, L. (2018). Age differences in memory for meaningful and arbitrary associations: A memory retrieval account. *Psychology and Aging*, *33*, 74–81.
- Amer, T., Giovanello, K. S., Nichol, D. R., Hasher, L., & Grady, C. L. (2019). Neural correlates of enhanced memory for meaningful associations with age. *Cerebral Cortex*. Advance online publication. doi:10.1093/cercor/bhy334
- Amer, T., & Hasher, L. (2014). Conceptual processing of distractors by older but not younger adults. *Psychological Science*, 25, 2252–2258.
- Amodio, D. M., & Frith, C. D. (2006). Meeting of minds: The medial frontal cortex and social cognition. *Nature Reviews Neuroscience*, 7, 268–277.
- Anderson, N. D., & Craik, F. I. (2017). 50 years of cognitive aging theory. The Journals of Gerontology, Series B: Psychological Sciences and Social Sciences, 72, 1–6.
- Andrews-Hanna, J. R., Reidler, J. S., Sepulcre, J., Poulin, R., & Buckner, R. L. (2010). Functional-anatomic fractionation of the brain's default network. *Neuron*, 65, 550–562.
- Andrews-Hanna, J. R., Smallwood, J., & Spreng, R. N. (2014). The default network and self-generated thought: Component processes, dynamic control, and clinical relevance. *Annals of the New York Academy of Sciences*, 1316, 29–52.
- Andrews-Hanna, J. R., Snyder, A. Z., Vincent, J. L., Lustig, C., Head, D., Raichle, M. E., & Buckner, R. L. (2007). Disruption of large-scale brain systems in advanced aging. *Neuron*, 56, 924–935.
- Baird, B., Smallwood, J., Gorgolewski, K. J., & Margulies, D. S. (2013). Medial and lateral networks in anterior prefrontal cortex support meta cognitive ability for memory and perception. *Journal of Neuroscience*, *33*, 16657–16665.
- Baltes, P. B., & Staudinger, U. M. (2000). Wisdom. A metaheuristic (pragmatic) to orchestrate mind and virtue toward excellence. *American Psychologist*, 55, 122–136.
- Bar, M., Aminoff, E., Mason, M., & Fenske, M. (2007). The units of thought. *Hippocampus*, 17, 420–428.
- Baror, S., & Bar, M. (2016). Associative activation and its relation to exploration and exploitation in the brain. *Psychological Science*, *27*, 776–789.
- Binder, J. R., Desai, R. H., Graves, W. W., & Conant, L. L. (2009). Where is the semantic system? A critical review and meta-analysis of 120 functional neuroimaging studies. *Cerebral Cortex*, 19, 2767–2796.
- Blanco, N. J., Love, B. C., Ramscar, M., Otto, A. R., Smayda, K., & Maddox, W. T. (2016). Exploratory decision-making as a function of lifelong experience, not cognitive decline. *Journal of Experimental Psychology: General*, 145, 284–297.
- Brainerd, C. J., & Reyna, V. F. (2015). Fuzzy-trace theory and lifespan cognitive development. *Developmental Review*, 38, 89–121.
- Brainerd, C. J., Reyna, V. F., & Howe, M. L. (2009). Trichotomous processes in early memory development, aging, and neurocognitive impairment: A unified theory. *Psychological Review*, 116, 783–832.
- Brashier, N. M., Umanath, S., Cabeza, R., & Marsh, E. J. (2017). Competing cues: Older adults rely on knowledge in the face of fluency. *Psychology and Aging*, *32*, 331–337.

Buckner, R. L. (2004). Memory and executive function in aging and AD: Multiple factors that cause decline and reserve factors that compensate. *Neuron*, 44, 195–208.

- Buckner, R. L., Andrews-Hanna, J. R., & Schacter, D. L. (2008). The brain's default network: Anatomy, function, and relevance to disease. *Annals of the New York Academy of Sciences*, 1124, 1–38.
- Cabeza, R. (2002). Hemispheric asymmetry reduction in older adults: The HAROLD model. *Psychology and Aging*, 17, 85–100.
- Carpenter, P. A., Just, M. A., & Shell, P. (1990). What one intelligence test measures: A theoretical account of the processing in the Raven Progressive Matrices Test. *Psychological Review*, 97, 404–431.
- Carstensen, L. L. (1992). Social and emotional patterns in adulthood: Support for socioemotional selectivity theory. *Psychology and Aging*, *7*, 331–338.
- Carstensen, L. L., Fung, H. H., & Charles, S. T. (2003). Socioemotional selectivity theory and the regulation of emotion in the second half of life. *Motivation and Emotion*, 27, 103–123.
- Castel, A. D. (2005). Memory for grocery prices in younger and older adults: The role of schematic support. *Psychology and Aging*, *20*, 718–721.
- Cattell, R. B. (1971). *Abilities: Their structure, growth, and action*. New York, NY: Houghton Mifflin.
- Chan, M. Y., Park, D. C., Savalia, N. K., Petersen, S. E., & Wig, G. S. (2014). Decreased segregation of brain systems across the healthy adult lifespan. *Proceedings of the National Academy of Sciences, USA, 111*, E4997–E5006.
- Cohen-Shikora, E. R., Diede, N. T., & Bugg, J. M. (2018). The flexibility of cognitive control: Age equivalence with experience guiding the way. *Psychology and Aging*, 33, 924–939.
- Craik, F. I., & Bialystok, E. (2006). Cognition through the lifespan: Mechanisms of change. *Trends in Cognitive Sciences*, *10*, 131–138.
- Craik, F. I. M., & Jennings, J. M. (1992). Human memory. In F. I. M. Craik & T. A. Salthouse (Eds.), *The handbook of aging and cognition* (pp. 51–110). Hillsdale, NJ: Erlbaum.
- Dalla Barba, G., Attali, E., & La Corte, V. (2010). Confabulation in healthy aging is related to interference of overlearned, semantically similar information on episodic memory recall. *Journal of Clinical and Experimental Neuropsychology*, *32*, 655–660.
- Damoiseaux, J. S. (2017). Effects of aging on functional and structural brain connectivity. *NeuroImage*, 160, 32–40.
- Davis, S. W., Dennis, N. A., Daselaar, S. M., Fleck, M. S., & Cabeza, R. (2008). Que pasa? The posterior-anterior shift in aging. *Cerebral Cortex*, *18*, 1201–1209.
- Deese, J. (1959). On the prediction of occurrence of particular verbal intrusions in immediate recall. *Journal of Experimental Psychology*, 58, 17–22.
- Duncan, J. (2010). The multiple-demand (MD)system of the primate brain: Mental programs for intelligent behaviour. *Trends in Cognitive Sciences*, *14*, 172–179.
- Duzel, E., Bunzeck, N., Guitart-Masip, M., & Duzel, S. (2010). Novelty-related motivation of anticipation and exploration by dopamine (nomad): Implications for healthy aging. *Neuroscience & Biobehavioral Reviews*, *34*, 660–669.

Forbes, C. E., & Grafman, J. (2010). The role of the human prefrontal cortex in social cognition and moral judgment. *Annual Review of Neuroscience*, *33*, 299–324.

- Gazzaley, A., Cooney, J. W., Rissman, J., & D'Esposito, M. (2005). Top-down suppression deficit underlies working memory impairment in normal aging. *Nature Neuro*science, 8, 1298–1300.
- Geerligs, L., Renken, R. J., Saliasi, E., Maurits, N. M., & Lorist, M. M. (2015). A brain-wide study of age-related changes in functional connectivity. *Cerebral Cortex*, 25, 1987–1999.
- Grady, C. (2012). The cognitive neuroscience of ageing. *Nature Reviews Neuroscience*, *13*, 491–505.
- Grady, C., Sarraf, S., Saverino, C., & Campbell, K. (2016).
  Age differences in the functional interactions among the default, frontoparietal control, and dorsal attention networks. *Neurobiology of Aging*, 41, 159–172.
- Hafkemeijer, A., van der Grond, J., & Rombouts, S. A. (2012). Imaging the default mode network in aging and dementia. Biochimica et Biophysica Acta: Protein Structure and Molecular Enzymology, 1822, 431–441.
- Hasher, L., & Zacks, R. T. (1988). Working memory, comprehension, and aging: A review and a new view. In G. H. Brower (Ed.), *The psychology of learning and motivation* (pp. 193–225). San Diego, CA: Academic Press.
- Hedden, T., Lautenschlager, G., & Park, D. C. (2005). Contributions of processing ability and knowledge to verbal memory tasks across the adult life-span. *Quarterly Journal of Experimental Psychology A*, 58, 169–190.
- Hills, T. T., Mata, R., Wilke, A., & Samanez -Larkin, G. R. (2013). Mechanisms of age-related decline in memory search across the adult life span. *Developmental Psychology*, 49, 2396–2404.
- Hills, T. T., Todd, P. M., Lazer, D., Redish, A. D., Couzin, I. D., & Cognitive Search Research Group. (2015). Exploration versus exploitation in space, mind, and society. *Trends in Cognitive Sciences*, 19, 46–54.
- Hoffman, P. (2018). Divergent effects of healthy ageing on semantic knowledge and control: Evidence from novel comparisons with semantically impaired patients. *Journal of Neuropsychology*. Advance online publication. doi:10.1111/jnp.12159
- Hoffman, P., & Morcom, A. M. (2018). Age-related changes in the neural networks supporting semantic cognition: A meta-analysis of 47 functional neuroimaging studies. *Neuroscience & Biobehavioral Reviews*, 84, 134–150.
- Jefferies, E. (2013). The neural basis of semantic cognition: Converging evidence from neuropsychology, neuroimaging and TMS. Cortex, 49, 611–625.
- Keller, J. B., Hedden, T., Thompson, T. W., Anteraper, S. A., Gabrieli, J. D., & Whitfield- Gabrieli, S. (2015). Restingstate anticorrelations between medial and lateral prefrontal cortex: Association with working memory, aging, and individual differences. *Cortex*, 64, 271–280.
- Konishi, M., McLaren, D. G., Engen, H., & Smallwood, J. (2015). Shaped by the past: The default mode network supports cognition that is independent of immediate perceptual input. *PLOS ONE*, 10(6), Article e0132209. doi:10.1371/journal.pone.0132209

- Koutstaal, W., Reddy, C., Jackson, E. M., Prince, S., Cendan, D. L., & Schacter, D. L. (2003). False recognition of abstract versus common objects in older and younger adults: Testing the semantic categorization account. *Journal of Experimental Psychology: Learning, Memory, and Cognition*, 29, 499–510.
- Koutstaal, W., Schacter, D. L., & Bernner, C. (2001). Dual task demands and gist-based false recognition of pictures in younger and older adults. *Journal of Memory and Language*, 44, 399–426.
- Leveroni, C. L., Seidenberg, M., Mayer, A. R., Mead, L. A., Binder, J. R., & Rao, S. M. (2000). Neural systems underlying the recognition of familiar and newly learned faces. *Journal of Neuroscience*, 20, 878–886.
- Levine, B., Svoboda, E., Hay, J. F., Winocur, G., & Moscovitch, M. (2002). Aging and autobiographical memory: Dissociating episodic from semantic retrieval. *Psychology and Aging*, *17*, 677–689.
- Li, S. C., Lindenberger, U., & Sikstrom, S. (2001). Aging cognition: From neuromodulation to representation. *Trends in Cognitive Sciences*, 5, 479–486.
- Li, S. C., Naveh-Benjamin, M., & Lindenberger, U. (2005). Aging neuromodulation impairs associative binding: A neurocomputational account. *Psychological Science*, 16, 445–450.
- Li, S. C., & Rieckmann, A. (2014). Neuromodulation and aging: Implications of aging neuronal gain control on cognition. *Current Opinion in Neurobiology*, 29, 148–158.
- Li, Y., Baldassi, M., Johnson, E. J., & Weber, E. U. (2013). Complementary cognitive capabilities, economic decision making, and aging. *Psychology and Aging*, 28, 595–613.
- Li, Y., Gao, J., Enkavi, A. Z., Zaval, L., Weber, E. U., & Johnson, E. J. (2015). Sound credit scores and financial decisions despite cognitive aging. *Proceedings of the National Academy of Sciences, USA*, 112, 65–69.
- Lockenhoff, C. E. (2011). Age, time, and decision making: From processing speed to global time horizons. *Annals of the New York Academy of Sciences*, 1235, 44–56.
- Lustig, C., & Buckner, R. L. (2004). Preserved neural correlates of priming in old age and dementia. *Neuron*, 42, 865–875.
- MacKay, D. G., Abrams, L., & Pedroza, M. J. (1999). Aging on the input versus output side: Theoretical implications of age-linked asymmetries between detecting versus retrieving orthographic information. *Psychology and Aging*, *14*, 3–17.
- Matzen, L. E., & Benjamin, A. S. (2013). Older and wiser: Older adults' episodic word memory benefits from sentence study contexts. *Psychology and Aging*, 28, 754–767.
- Meyer, M. L., Spunt, R. P., Berkman, E. T., Taylor, S. E., & Lieberman, M. D. (2012). Evidence for social working memory from a parametric functional MRI study. *Proceedings of the National Academy of Sciences, USA*, 109, 1883–1888.
- Musielak, C., Giraudeau, C., Chasseigne, G., & Mullet, E. (2014). To what extent does the existence of functional relations in a learning setting change the pattern of differences between younger and older adults' performances? *Experimental Aging Research*, 40, 455–476.

- Naveh-Benjamin, M., Craik, F. I., Guez, J., & Kreuger, S. (2005). Divided attention in younger and older adults: Effects of strategy and relatedness on memory performance and secondary task costs. *Journal of Experimental Psychology: Learning, Memory, and Cognition*, *31*, 520–537.
- Ng, K. K., Lo, J. C., Lim, J. K., Chee, M. W., & Zhou, J. (2016). Reduced functional segregation between the default mode network and the executive control network in healthy older adults: A longitudinal study. *NeuroImage*, 133, 321–330.
- Park, D. C., Polk, T. A., Mikels, J. A., Taylor, S. F., & Marshuetz, C. (2001). Cerebral aging: Integration of brain and behavioral models of cognitive function. *Dialogues in Clinical Neuroscience*, 3, 151–165.
- Park, D. C., & Reuter-Lorenz, P. (2009). The adaptive brain: Aging and neurocognitive scaffolding. *Annual Review of Psychology*, 60, 173–196.
- Payer, D., Marshuetz, C., Sutton, B., Hebrank, A., Welsh, R. C., & Park, D. C. (2006). Decreased neural specialization in old adults on a working memory task. *NeuroReport*, 17, 487–491.
- Persson, J., Lustig, C., Nelson, J. K., & Reuter-Lorenz, P. A. (2007). Age differences in deactivation: A link to cognitive control? *Journal of Cognitive Neuroscience*, 19, 1021–1032.
- Piguet, O., Double, K. L., Kril, J. J., Harasty, J., Macdonald, V., McRitchie, D. A., & Halliday, G. M. (2009). White matter loss in healthy ageing: A postmortem analysis. *Neurobiology of Aging*, 30, 1288–1295.
- Raichle, M. E., MacLeod, A. M., Snyder, A. Z., Powers, W. J., Gusnard, D. A., & Shulman, G. L. (2001). A default mode of brain function. *Proceedings of the National Academy* of Sciences, USA, 98, 676–682.
- Ralph, M. A., Jefferies, E., Patterson, K., & Rogers, T. T. (2017). The neural and computational bases of semantic cognition. *Nature Reviews Neuroscience*, 18, 42–55.
- Ramscar, M., Hendrix, P., Shaoul, C., Milin, P., & Baayen, H. (2014). The myth of cognitive decline: Non-linear dynamics of lifelong learning. *Topics in Cognitive Science*, 6, 5–42.
- Raz, N., Lindenberger, U., Rodrigue, K. M., Kennedy, K. M., Head, D., Williamson, A., . . . Acker, J. D. (2005). Regional brain changes in aging healthy adults: General trends, individual differences and modifiers. *Cerebral Cortex*, 15, 1676–1689.
- Reuter-Lorenz, P. A., & Cappell, K. A. (2008). Neurocognitive aging and the compensation hypothesis. *Current Directions in Psychological Science*, 17, 177–182.
- Reyna, V. F. (2012). A new intuitionism: Meaning, memory, and development in fuzzy-trace theory. *Judgment and Decision Making*, 7, 332–359.
- Reyna, V. F., & Brainerd, C. J. (1995). Fuzzy-trace theory: An interim synthesis. *Learning and Individual Differences*, 7, 1–75.
- Reyna, V. F., & Brainerd, C. J. (2011). Dual processes in decision making and developmental neuroscience: A fuzzy-trace model. *Developmental Review*, *31*, 180–206.
- Rieck, J. R., Rodrigue, K. M., Boylan, M. A., & Kennedy, K. M. (2017). Age-related reduction of BOLD modulation to cognitive difficulty predicts poorer task accuracy and poorer fluid reasoning ability. *NeuroImage*, 147, 262–271.

- Roediger, H. L. I., & McDermott, K. B. (1995). Creating false memories: Remembering words not presented in lists *Journal of Experimental Psychology: Learning, Memory, and Cognition*, *21*, 803–814.
- Ruch, F. L. (1934). The differentiative effects of age upon human learning. *Journal of General Psychology*, 11, 261–286.
- Salthouse, T. A. (1996). The processing-speed theory of adult age differences in cognition. *Psychological Review*, *103*, 403–428.
- Samanez-Larkin, G. R. (2013). Financial decision making and the aging brain. *APS Observer*, 26(5), 30–33.
- Samanez-Larkin, G. R., & Knutson, A. (2014). Reward processing and risky decision making in the aging brain. In V. Reyna & V. Zayas (Eds.), *The neuroscience of risky decision making* (pp. 123–142). Washington, DC: American Psychological Association.
- Samanez-Larkin, G. R., & Knutson, B. (2015). Decision making in the ageing brain: Changes in affective and motivational circuits. *Nature Reviews Neuroscience*, *16*, 278–289.
- Schacter, D. L., Koutstaal, W., Johnson, M. K., Gross, M. S., & Angell, K. E. (1997). False recollection induced by photographs: A comparison of older and younger adults. *Psychology and Aging*, 12, 203–215.
- Schacter, D. L., Koutstaal, W., & Norman, K. A. (1997). False memories and aging. *Trends in Cognitive Sciences*, 1, 229–236
- Schmitz, T. W., Dixon, M. L., Anderson, A. K., & De Rosa, E. (2014). Distinguishing attentional gain and tuning in young and older adults. *Neurobiology of Aging*, *35*, 2514–2525.
- Schwartenbeck, P., Fitzgerald, T., Dolan, R. J., & Friston, K. (2013). Exploration, novelty, surprise, and free energy minimization. *Frontiers in Psychology*, *4*, Article 710. doi:10.3389/fpsyg.2013.00710
- Shallice, T. (1982). Specific impairments of planning. *Philosophical Transactions of the Royal Society B: Biological Sciences*, 298, 199–209.
- Shallice, T., & Burgess, P. (1996). The domain of supervisory processes and temporal organization of behaviour. *Philosophical Transactions of the Royal Society B: Biological Sciences*, *351*, 1405–1411; discussion 1411–1412.
- Sheldon, S., McAndrews, M. P., & Moscovitch, M. (2011). Episodic memory processes mediated by the medial temporal lobes contribute to open-ended problem solving. *Neuropsychologia*, 49, 2439–2447.
- Snowdon, D. A. (1997). Aging and Alzheimer's disease: Lessons from the Nun Study. Gerontologist, 37, 150–156.
- Spaniol, J., & Bayen, U. J. (2002). When is schematic knowledge used in source monitoring? *Journal of Experimental Psychology: Learning, Memory, and Cognition*, 28, 631–651.
- Spreng, R. N. (2012). The fallacy of a "task-negative" network. *Frontiers in Psychology*, *3*, Article 145. doi:10.3389/fpsyg.2012.00145
- Spreng, R. N., DuPre, E., Selarka, D., Garcia, J., Gojkovic, S., Mildner, J., . . . Turner, G. R. (2014). Goal-congruent default network activity facilitates cognitive control. *Journal of Neuroscience*, 34, 14108–14114.
- Spreng, R. N., Gerlach, K. D., Turner, G. R., & Schacter, D. L. (2015). Autobiographical planning and the brain:

Activation and its modulation by qualitative features. *Journal of Cognitive Neuroscience*, *27*, 2147–2157.

- Spreng, R. N., Lockrow, A. W., DuPre, E., Setton, R., Spreng, K. A. P., & Turner, G. R. (2018). Semanticized autobiographical memory and the default–executive coupling hypothesis of aging. *Neuropsychologia*, *110*, 37–43.
- Spreng, R. N., & Schacter, D. L. (2012). Default network modulation and large-scale network interactivity in healthy young and old adults. *Cerebral Cortex*, 22, 2610–2621.
- Spreng, R. N., Stevens, W. D., Chamberlain, J., Gilmore, A. W., & Schacter, D. L. (2010). Default network activity, coupled with the frontoparietal control network, supports goaldirected cognition. *NeuroImage*, 53, 303–317.
- Spreng, R. N., Stevens, W. D., Viviano, J. D., & Schacter, D. L. (2016). Attenuated anticorrelation between the default and dorsal attention networks with aging: Evidence from task and rest. *Neurobiology of Aging*, *45*, 149–160.
- Spreng, R. N., & Turner, G. R. (2013). Structural covariance of the default network in healthy and pathological aging. *Journal of Neuroscience*, *33*, 15226–15234.
- Spreng, R. N., & Turner, G. R. (2019). Structure and function of the aging brain. In G. R. Samanez-Larkin (Ed.), *The aging brain: Functional adaptation across adulthood.* Washington, DC: American Psychological Association.
- Sternberg, R. J. (1998). A balance theory of wisdom. *Review of General Psychology*, 2, 347–365.
- Stevens, W. D., & Spreng, R. N. (2014). Resting-state functional connectivity MRI reveals active processes central to cognition. Wiley Interdisciplinary Reviews: Cognitive Science, 5, 233–245.
- Stokes, M. G., Kusunoki, M., Sigala, N., Nili, H., Gaffan, D., & Duncan, J. (2013). Dynamic coding for cognitive control in prefrontal cortex. *Neuron*, 78, 364–375.

- Stuss, D. T., & Levine, B. (2002). Adult clinical neuropsychology: Lessons from studies of the frontal lobes. *Annual Review of Psychology*, *53*, 401–433.
- Turner, G. R., & D'Esposito, M. (2010). Functional neuroimaging in aging. In J. Knoefel & M. Albert (Eds.), *Clinical neurology of aging* (3rd ed., pp. 105–112). New York, NY: Oxford University Press.
- Turner, G. R., & Spreng, R. N. (2015). Prefrontal engagement and reduced default network suppression cooccur and are dynamically coupled in older adults: The default-executive coupling hypothesis of aging. *Journal of Cognitive Neuroscience*, 27, 2462–2476.
- Umanath, S., & Marsh, E. J. (2012). Aging and the memorial consequences of catching contradictions with prior knowledge. *Psychology and Aging*, *27*, 1033–1038.
- Umanath, S., & Marsh, E. J. (2014). Understanding how prior knowledge influences memory in older adults. *Perspectives on Psychological Science*, *9*, 408–426.
- Vandermorris, S., Sheldon, S., Winocur, G., & Moscovitch, M. (2013). Differential contributions of executive and episodic memory functions to problem solving in younger and older adults. *Journal of the International Neuro-psychological Society*, 19, 1087–1096.
- Verhaeghen, P. (2003). Aging and vocabulary scores: A metaanalysis. *Psychology and Aging*, *18*, 332–339.
- Verhaeghen, P., & Cerella, J. (2002). Aging, executive control, and attention: A review of meta-analyses. *Neuroscience & Biobehavioral Reviews*, 26, 849–857.
- Yarchoan, M., Xie, S. X., Kling, M. A., Toledo, J. B., Wolk, D. A., Lee, E. B., . . . Arnold, S. E. (2012). Cerebrovascular atherosclerosis correlates with Alzheimer pathology in neurodegenerative dementias. *Brain*, *135*(Pt. 12), 3749–3756.