Research report

Resting-state anticorrelations between medial and lateral prefrontal cortex: Association with working memory, aging, and individual differences

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ABSTRACT

We examined how variation in working memory (WM) capacity due to aging or individual differences among young adults is associated with intrinsic or resting-state anticorrelations, particularly between (1) the medial prefrontal cortex (MPFC), a component of the default-mode network (DMN) that typically decreases in activation during external, attention-demanding tasks, and (2) the dorsolateral prefrontal cortex (DLPFC), a component of the fronto-parietal control network that supports executive functions and WM and typically increases in activation during attention-demanding tasks. We compared the magnitudes of MPFC-DLPFC anticorrelations between healthy younger and older participants (Experiment 1) and related the magnitudes of these anticorrelations to individual differences on two behavioral measures of WM capacity in two independent groups of young adults (Experiments 1 and 2). Relative to younger adults, older adults exhibited reductions in WM capacity and in MPFC-DLPFC anticorrelations. Within younger adults, greater MPFC-DLPFC anticorrelation at rest correlated with greater WM capacity. These findings show that variation in MPFC-DLPFC anticorrelations, whether related to aging or to individual differences, may reflect an intrinsic functional brain architecture supportive of WM capacity.

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1. Introduction

Working memory (WM) capacity, defined as the amount of goal-relevant information that can be both maintained and manipulated, declines with age (Craik, Morris, & Gick, 1990) and varies considerably among individuals (Engle, 2002). In contrast to measures of simple short-term maintenance of information (e.g., digit span), greater WM capacity is associated with superior performance in a broad range of high-level cognitive domains, including reading comprehension, problem solving, and inhibitory control (Conway, Kane, & Engle, 2003). WM capacity is thought to reflect central executive capability (Baddeley, 1992; Engle, 2002), and to depend on dorsolateral prefrontal cortex (DLPFC), parietal cortex, anterior cingulate cortex, and the basal ganglia (D’Esposito, 2007; D’Esposito, Postle, Ballard, & Lease, 1999; Frank, Loughry, & O’Reilly, 2001; Levy & Goldman-Rakic, 2000). Here, we asked whether a relationship exists between variation in WM capacity, due to aging or across younger individuals, and the intrinsic functional architecture of the human brain as measured by resting-state functional connectivity.

Spontaneous fluctuations in functionally related brain regions are correlated with each other in the absence of external stimuli, and the patterns of these correlations have been thought to reveal intrinsic relations of brain regions (Beckmann, DeLuca, Devlin, & Smith, 2005; Biswal, Yetkin, Haughton, & Hyde, 1995; Greicius, Krasnow, Reiss, & Menon, 2003). During rest, in young adults, there are strong correlations between components of the default-mode network (DMN), brain regions that are commonly deactivated during external or attention-demanding tasks involving mental control (Fox et al., 2005; Fransson, 2005; Greicius et al., 2003; Raichle et al., 2001). Anatomically, the DMN includes medial prefrontal cortex (MPFC), posterior cingulate cortex (PCC), left and right lateral parietal (LLP and RLP) cortices, and bilateral medial temporal lobe (MTL) regions (Buckner, Andrews-Hanna, & Schacter, 2008).

Resting-state correlations among the components of the DMN appear to be significantly reduced in age-associated pathologies (Greicius, Srivastava, Reiss, & Menon, 2004; Hedden et al., 2009) and in typically aging older adults (Andrews-Hanna et al., 2007; Balsters et al., 2013; Damoiseaux et al., 2008; Grady et al., 2010; Mowinckel, Espeseth, & Westlye, 2012; Sala-Llonch et al., 2012; Sambataro et al., 2010). This may be due, in part, to the particular vulnerability of long-range DMN functional connections to the effects of normal aging (Allen et al., 2011; Andrews-Hanna et al., 2007; Fillippini et al., 2012; Hafkemeijer, van der Grond, & Rombouts, 2012; Tomasi & Volkow, 2012) or a consequence of more motion-related artificial time points in elderly participants (Power, Barnes, Snyder, Schlaggar, & Petersen, 2012; Van Dijk, Sabuncu, & Buckner, 2012). Although widespread reductions in resting functional connectivity are observed with advancing age, some studies also report localized increases in resting-state functional connectivity. Older adults, relative to younger adults, have shown increased frontal-lobe coherence (Fillippini et al., 2012) and increased functional connectivity within frontoparietal cortical regions (Mowinckel et al., 2012).

Networks in the brain appear to have an intrinsic organization such that different networks may exhibit negative functional connectivity, or are anticorrelated with one another at rest. In young adults, components of the DMN are negatively correlated with brain networks comprised of regions commonly activated for external tasks that demand attention and mental control, including the DLPFC (Fox et al., 2005; Fransson, 2005). Evaluation of negatively correlated networks has proven controversial due to global signal regression, a method used commonly to mitigate physiological noise in resting-state functional imaging studies. Global signal regression is known to mathematically generate anticorrelations (Murphy, Birn, Handwerker, Jones, & Bandettini, 2009; Saad et al., 2012). Given these issues, valid analysis of negatively correlated networks has developed into a topic of particular interest in the field (Chang & Glover, 2009; Fox et al., 2005; Hampson, Driesen, Roth, Gore, & Constable, 2010; Saad et al., 2012; Van Dijk et al., 2010; Weissenbacher et al., 2009). With the caveat that prior studies of the influence of age on anticorrelations have employed global signal regression, there is evidence that healthy aging is also characterized by reduced negative correlations at rest between the DMN and cortical regions commonly recruited during attention-demanding tasks (Wu et al., 2011).

Variation in DMN connectivity has been associated with variation in executive functions and WM capacity. Among older adults, reduced MPFC-PCC connectivity correlated with worse performance on executive-function and other cognitive measures (Andrews-Hanna et al., 2007) and reduced connectivity in a DMN network dominated by the MPFC correlated with worse performance on a trail-making test (Damoiseaux et al., 2008). Neither study reported a correlation between these brain measures and variation among young adults, because that was either not examined (Andrews-Hanna et al., 2007) or was not significant in 10 participants (Damoiseaux et al., 2008). For young adults, there is a report of a positive correlation between magnitude of MPFC-DLPFC anticorrelation and WM capacity, as measured by an n-back task (Hampson et al., 2010). The relation between reduced MPFC-DLPFC resting-state anticorrelation and reduced WM capacity is consistent with findings from patients with schizophrenia (Whitfield-Gabrieli et al., 2009).

Although variability in resting-state functional connectivity has been associated with variation in WM in relation to aging and to individual differences among young adults, there are two major gaps in the current understanding of that association. First, studies of aging have implicated positive correlations with the MPFC as being related to age-associated reduction in WM capacity, whereas the one study of variation among young adults has, instead, implicated negative correlations with MPFC. This leaves open the question about whether age-related changes in WM and individual differences among young adults in WM capacity are associated with shared or distinct variations in intrinsic functional connectivity (no one study has discovered such common variation in both younger and older adults). Second, the above functional connectivity findings were reported before it was well understood that greater movement in older than younger adults can produce artifactual results (Power et al., 2012; Van Dijk et al., 2012) or that global signal regression can mathematically generate anticorrelations (Murphy et al., 2009; Saad et al.,...
Therefore, it is unknown whether the prior findings would hold when methodological improvements were implemented. Here, we explored whether there exists shared or distinct characteristics of intrinsic brain function for age-related declines in WM capacity and for individual differences among young adults in WM capacity. We focused on MPFC positive and negative functional connectivity because bi-directional correlations of the MPFC with different regions have been implicated across studies of aging or of individual differences among young adults in relation to executive functions and WM capacity (Andrews-Hanna et al., 2007; Damoiseaux et al., 2008; Hampson et al., 2010). We examined the relation of MPFC-DLPFC anticorrelations and MPFC-PCC positive correlations to WM capacity (Experiment 1) in 27 younger and 27 older healthy adults with capacity measured by the Letter-Number Sequencing subtest from the Wechsler Adult Intelligence Scale (WAIS-III), and in 70 younger adults (Experiment 2) with a composite measure of Operation and Reading Span tests (Turner & Engle, 1989; Unsworth, Heitz, Schrock, & Engle, 2005). In both experiments, we implemented methods that minimize the influence of motion artifacts and physiological noise and allow for valid interpretations of negative correlations (Behzadi, Restom, Liu, & Liu, 2007; Chai, Castañón, Ongür, & Whitfield-Gabrieli, 2012; Whitfield-Gabrieli & Nieto-Castanon, 2012).

2. Materials and methods

2.1. Experiment 1

2.1.1. Participants

Participants were 27 older adults (15 women) between 65 and 89 years of age ($M = 75.7$ years, $SD = 6.7$) and 27 younger adults (15 women) between 20 and 33 years of age ($M = 24.8$, $SD = 3.4$). Written informed consent for participation in the study was obtained from all participants and approved by the MIT Institutional Review Board. All participants were healthy, right-handed individuals (Oldfield, 1971) from the Boston metropolitan area who satisfied the following criteria: native English speakers; no contraindications to MRI; and absence of neurological or psychiatric impairments or associated medications. All participants had normal or corrected-to-normal vision. No participant exhibited evidence of mild cognitive impairment or dementia; participants were excluded if they scored $<27$ on the Mini-Mental State Examination (Folstein & Folstein, 1975).

2.1.2. Neuropsychological and demographic measures

The Letter-Number Sequencing subtest from the Wechsler Adult Intelligence Scale (WAIS-III) was used as the measure of WM capacity. Participants were read a combination of numbers and letters, and then asked to recall first the numbers in ascending order and then the letters in alphabetical order. The score was the maximum number of items reordered and recalled correctly from WM (Wechsler, 2002). Two measures were used to assess comparability of the age groups. The American version of the National Reading Test (AMNART) (Grober & Sliwinski, 1991) was used to estimate crystallized IQ. Socioeconomic status (SES) was measured with the Hollingshead SES scale, which separately ranks an individual’s educational and occupational attainment on scales ranging from 1 to 7. A weighted score was computed by multiplying the educational score by 4 and the occupational score by 7 and summing the 2 scores (Hollingshead, 1957). Lower scores indicate higher SES. Because the majority of younger participants had not yet completed their educations, we compared the older group to the SES scores for the parents of the younger group.

2.1.3. MRI data acquisition

Functional magnetic resonance imaging (fMRI) data were acquired using a 3-Tesla Siemens Tim Trio scanner (Siemens, Erlangen, Germany) paired with a 12-channel phased-array whole-head coil. Head motion was restrained with foam pillows and extendable padded head clamps 3D T1-weighted magnetization prepared rapid acquisition gradient echo (MP-RAGE) anatomical images were collected with the following parameters: time repetition (TR) = 2530 msec, time echo (TE) = 3.39 msec, flip angle (FA) = 7°, 1.33 $\times$ 1.33 mm resolution, 2$\times$ acceleration. Functional T2*-weighted images were acquired using a gradient -echo planar pulse sequence sensitive to bold oxygenation level-dependent (BOLD) contrast (Kwong et al., 1992; Ogawa et al., 1992) with the following parameters: TR = 2000 msec, TE = 30 msec, FA = 90°, 3.0 mm isotropic resolution. Thirty-six transverse slices covered the whole brainand were acquired in an interleaved fashion. Functional data were acquired while the participant was instructed to rest with eyes open for a period of 5 min consisting of 150 volumes. To allow for T1-equilibration effects, 4 dummy volumes were discarded prior to acquisition. Online prospective acquisition correction (PACE) was applied to the EPI sequence.

2.1.4. Resting-state preprocessing

Resting-state fMRI data were first preprocessed in SPM5 (Wellcome Department of Imaging Neuroscience, London, UK; [http://www.fil.ion.ucl.ac.uk/spm/spm5.html]). Images were realigned (motion corrected), spatially normalized to the Montreal Neurological Institute (MNI) stereotactic space, and smoothed with a 6 mm kernel. Quality assurance was performed on the functional time series in order to detect outliers in the motion and global signal intensity using the in-house software art (http://www.nitrc.org/projects/artifact_detect). From each participant, an image was identified as an outlier if composite movement from a preceding image exceeded 0.5 mm, or if the global mean intensity was greater than 3 standard deviations from the mean image intensity for the run in which it was collected. This composite motion measure was defined by the art tool. By default, art converts the 6 rotation/translation head motion parameters into another set of 6 parameters characterizing the trajectories of 6 points located on the center of each of the faces of a bounding box around the brain. It then computes the maximum scan-to-scan movement of any of these points as the single ‘composite’ scan-to-scan movement measure, which is thresholded to determine outliers. Identified outliers were included as nuisance parameters, as one regressor per outlier, with the first level general linear models.
2.1.5. Functional connectivity analysis

Functional connectivity analysis was performed with a seed-driven approach using the in-house, custom software Conn (http://www.nitrc.org/projects/conn; Whitfield-Gabrieli & Nieto-Castanon, 2012). The MPFC seed was defined a priori from the literature (Fox et al., 2005; Whitfield-Gabrieli et al., 2009) as 10 mm spheres around the coordinates for the MPFC (−1, 47, −4) in MNI space. Physiological and other spurious sources of noise were estimated using the aCompCor method (Behzadi et al., 2007; Chai et al. 2012; Whitfield-Gabrieli et al., 2009), and removed together with movement-related and artifactual covariates. The residual BOLD time-series was band-pass filtered (.008Hz–.08 Hz). Each participant's structural image was segmented into white matter, gray matter (GM), and cerebral spinal fluid (CSF) using SPM8. WM and CSF masks were eroded by one voxel to avoid partial volume effects with adjacent GM. The first 3 principal components of the signals from the eroded white matter and CSF noise ROIs were removed with regression.

First-level correlation maps were produced by extracting the residual BOLD time course from the MPFC seed and computing Pearson's correlation coefficients between that time course and the time courses of all other voxels in the brain. Correlation coefficients were Fisher transformed into ‘Z’ scores, which increases normality and allows for improved second-level General Linear Model analyses. All reported clusters were significant at an FDR cluster-corrected threshold of $p < 0.05$.

2.1.5.1. Within group analyses. MPFC seed-to-voxel group analyses were separately performed using one-sample t-tests for within the young cohort ($n = 27$), and the older cohort ($n = 27$).

2.1.5.2. Between group analyses (Older versus Younger). MPFC seed-to-voxel between group connectivity analyses were performed using two sample t-tests.

2.1.5.3. MPFC functional connectivity correlation with WM. We investigated the relationship of (a) the magnitude of resting-state correlations from the MPFC that were either positively correlated with the PCC or negatively correlated with the bilateral DLPFC regions and (b) the measure of WM performance assessed outside of the scanner (Letter-Number Sequencing task). We performed a one-sample t-test for the entire group ($n = 54$) and functionally defined (a) the left and right DLPFC clusters (within BA 46/9) that were significantly anticorrelated with the MPFC, and (b) the PCC cluster (BA 30/31) that was significantly positively correlated with the MPFC seed. We then extracted the mean $Z$-values from the bilateral DLPFC and PCC clusters for each participant of both cohorts and correlated those values with their WM capacities (as defined by the letter-number sequencing task) within both groups. Thus, the ROIs were unbiased because they were derived from all participants and independently from any behavioral measures.

2.1.6. Matched groups on motion artifacts

To ensure that between group results were not driven by age-related differences in motion artifacts, we performed additional between-group analyses of subgroups of older and younger adults who did not differ significantly on movement and other artifacts. Within each cohort, 5 participants were removed ($n = 44$) to create groups equated for artifacts. Two-sample t-tests were performed to directly compare the connectivity maps between older and younger adults in the movement and artifact-matched groups.

2.2. Experiment 2

2.2.1. Participants

Participants were 70 younger adults (39 women) between the ages of 18 and 29 years of age ($M = 21.6$ years, SD = $2.6$). Written informed consent was approved by the MIT Institutional Review Board. Participants were required to be adults between the ages of 18 and 45, right-handed, in good health, and not taking any drugs. They were recruited through web advertisements, physical flyers, and e-mail to the Northeastern and Tufts college mailing lists.

2.2.2. Neuropsychological and demographic measures

The automated Operation Span and Reading Span tasks were used as measures of complex WM capacity (Unsworth et al., 2005). For the Operation Span task, participants were presented with alternating letters and math equations, and asked to remember the letters while assessing whether each equation was valid. Set sizes ranged from 3-letters to 7-letters, with each set size presented for 3 trials over the course of the task, in a random order. At the end of each trial, participants reported the letters in the order they were presented. The dependent measure was the sum of all perfectly remembered letter sets. For the Reading Span task, participants were presented with alternating letters and sentences, and asked to remember the letters while assessing whether each sentence was sensible. Set sizes and scoring were identical to the automated Operation Span. Finally, Operation Span and Reading Span scores were summed to create a single measure estimating a participant’s complex WM capacity (composite score).

2.2.3. MRI data acquisition

Data were acquired on a 3T Tim Trio Siemens scanner using a 32-channel head coil. T1-weighted whole brain anatomical images (MPRAGE sequence, 256 x 256 voxels, 1 x 1.3-mm in-plane resolution, 1.3-mm slice thickness) were acquired. All participants underwent a resting functional MRI scan of 6 min with the instructions “keep your eyes closed and think of nothing in particular”. Resting scan images were obtained in 62 2-mm thick transverse slices, covering the entire brain (interleaved EPI sequence, T2*-weighted images; repetition time = 6 sec, echo time = 30 msec, flip angle = 90, 67 slices with 2 x 2 x 2 mm voxels). FACE was applied to the EPI sequence.

2.2.4. Data analysis

Resting-state fMRI data for Experiment 2 were first preprocessed in SPM8 (Wellcome Department of Imaging Neuroscience, London, UK; http://www.fil.ion.ucl.ac.uk/brain), using standard spatial preprocessing steps. Images were slice-
time corrected, realigned and resliced, normalized in MNI space and smoothed with a 4-mm kernel.

2.2.5. Functional connectivity analysis
The first four scans were excluded from analysis, as there were no dummy scans during initial acquisition. The preprocessing, artifact detection and rejection, aCompCor denoising to address physiological aliasing and subsequent seed driven functional connectivity analyses were identical to those in Experiment 1. Importantly, the left and right DLPFC clusters were defined from the entire group’s anticorrelation with the MPFC (independent of WM) and the mean DLPFC Z-values from the clusters were then correlated with the complex WM scores.

2.3. Results

2.3.1. Neuropsychological measures for Experiment 1
There was no significant difference between younger (M = 119.74, SD = 5.8) and older (M = 120.37, SD = 7.6) groups for AMNART scores ([t(52)] = .36, p = .73) or for the Hollingshead SES scale younger: M = 28.37, SD = 10.0; older: M = 31.96, SD = 11.6; [t(52)] = 1.22, p = .22. The younger group (M = 15.15, SD = 3.4) performed significantly better than the older group (M = 10.70, SD = 2.5) on the Letter-Number Sequencing Test ([t(52)] = 5.44, p < .001]. Analyses of behavioral measures were performed with two-tailed t-tests.

2.3.2. Artifact detection
Relative to younger adults (M = 1.1% of 150 time points, SD = 1.6%), older adults (M = 2.4% of 150 time points, SD = 3.0%) had significantly more artifacts (the union of motion and intensity outliers) ([t(52)] = 2.05, p < .05]. MPFC correlations and anticorrelations increased post artifact detection and rejection, most noticeably in the older adults.

In order to make certain that group differences between younger and older adults were not not driven by age-related differences in motion artifacts, we performed between group analyses on the groups who were matched on motion artifacts [after eliminating five participants from each cohort (n = 44)]. For these matched groups, there was no significant difference in motion and other artifacts ([t(42)] = .00, p = 1.00].

2.3.3. Group differences in intrinsic functional organization
Using a seed-to-voxel analysis, the positive correlation between the MPFC seed and the PCC was significantly reduced in older relative to younger adults, (Fig. 1A, middle column), and younger > older adults (right column). Top row depicts results from all participants. Second row depicts results from groups matched for motion artifacts. Results thresholded at p < .05, FDR cluster corrected in top two rows. The third row shows the same analyses at a more liberal threshold (p = .05, unc), and reveals that age-related differences occur only in frontal regions even at this threshold. (B) Resting-state anticorrelation for younger (gray) and older (black) adults between MPFC and bilateral DLPFCs; only younger adults exhibited significant anticorrelations in left and right DLPFC.

2.3.4. Correlations with WM performance
There was not a significant correlation between greater MPFC-PCC correlation and superior performance on the Letter-Number Sequencing task within either the older (r = .01, p = .95) or the younger group (r = .18, p = .38). Greater anticorrelation between the MPFC and the right DLPFC was significantly associated with better performance within the younger group (r = -.43, p < .05), but not within the older group
The lack of association within the older group may be attributable to a restricted range of variance in the greatly reduced anticorrelation between MPFC and right DLPFC. Correlations in the younger adults remained significant after removing a young adult with an apparent outlier value in the MPFC/DLPFC anticorrelations. The left DLPFC was not significantly correlated with WM performance within either group.

2.3.5. Results for Experiment 2
We replicated and extended our findings in Experiment 1 (that greater resting-state MPFC-DLPFC anticorrelation was associated with greater WM capacity among young adults) with a new independent group with a larger sample size (\( n = 70 \)). The primary analysis was performed with the composite score from two complex WM measures (i.e., Operation Span and Reading Span). Greater MPFC-left DLPFC anticorrelations were significantly correlated with composite WM scores (\( r = -.24, p = .04 \)) (Fig. 3).

3. Discussion
We found convergent evidence from aging and from individual differences among young adults of a relation between greater WM capacity and greater magnitude of MPFC-DLPFC anticorrelation. Older adults exhibited both reduced WM and reduced MPFC-DLPFC anticorrelation relative to younger adults. Furthermore, greater WM capacity was associated with greater MPFC-DLPFC anticorrelation in two independent cohorts of young adults (total \( n = 97 \)) with two different WM measures.

3.1. Age-related differences in WM capacity and MPFC-DLPFC anticorrelation
The behavioral findings were consistent with those generally observed in healthy aging (Hedden & Gabrieli, 2004, 2005). Younger and older participants scored similarly on the AMNART, a measure of vocabulary knowledge, consistent with evidence that crystallized knowledge or intelligence remains relatively intact during healthy aging (Park et al., 2002; Schaie, 1996). In contrast, older participants scored significantly less well on the measure of WM capacity, consistent with evidence that WM or fluid intelligence abilities decline in healthy aging (Park et al., 2002; Schaie, 1996). The validity of comparing these two groups of younger and older participants was supported by similar AMNART scores and similar SES status (with the use of parental SES for the younger adults who have often not completed education or reached final career and economic status).

There was also a significant age-related reduction in MPFC-DLPFC resting-state anticorrelation. This finding is consistent with a prior study that employed a potentially problematic method of global signal regression to examine anticorrelations (Wu et al., 2011). In addition, older adults, relative to younger adults, exhibited significant reductions in MPFC-PCC positive correlations. Reduced positive MPFC-PCC correlations are consistent with prior reports of reduced functional connectivity in normal aging (Andrews-Hanna et al., 2007; Damoiseaux et al., 2008). Aging appeared to have a more severe impact on the MPFC-DLPFC anticorrelation because that anticorrelation was statistically absent, whereas the reduced

\[ r = -.11, p = .51 \] (Fig. 2).

\[ r = -.43, p = .0241 \]

\[ r = -.11, p = .6016 \]
event-related activation design (e.g., Andrews-Hanna et al., present study, but perform correlations on the residuals of an studies do not measure resting-state correlations as in the neuropsychological tests, or other factors. For example, some approaches, sample characteristics, choice of either younger or older adults in Experiment 1. This difference across studies may be due to variations in analytic approaches, sample characteristics, choice of neuropsychological tests, or other factors. For example, some studies do not measure resting-state correlations as in the present study, but perform correlations on the residuals of an event-related activation design (e.g., Andrews-Hanna et al., 2007). Another study found a significant correlation between DMN anticorrelations during a 3-back task and WM performance on the 3-back task (Sala-Llonch et al., 2012). This relationship was not found in any of the less demanding n-back levels, but the anticorrelation during the interleaved fixation periods also correlated with WM performance during the 3-back condition (there was no report of the DMN anticorrelations during the pure rest condition and WM performance on the 3-back task). However, there is evidence that an active n-back task influences the rest periods within the task design, and therefore may be different from calculating the anticorrelations from pure rest (Pyka et al. 2009). In the present study, all resting-state measures were taken from pure resting-state scans, and are therefore not confounded with task performance.

The age-associated alteration in functional connectivity may be associated with age-related alteration in structural connectivity of white matter pathways as measured by diffusion tensor imaging (DTI). Older adults exhibit reduced integrity relative to younger people, especially in anterior regions near the DLPFC (e.g., Head et al., 2004; Pfefferbaum, Adalsteinsson, & Sullivan, 2005; Salat et al., 2005). Altered integrity of anterior white-matter pathways has been associated with age-related reductions in cognitive control and executive functions (Charlton et al., 2006; Deary et al., 2006; Grieve, Williams, Paul, Clark, & Gordon, 2007; O’Sullivan et al., 2001; Penke et al., 2010). Indeed, such region-specific alterations of white matter in typical aging have been related to individual differences in cognitive control among older adults, whereas cortical thickness and age-related alterations of temporal and parietal lobe white matter were unrelated (Ziegler et al., 2010).

Because many neural and psychological changes occur in concert with aging, it is difficult to make strong causal assertions between specific neural and specific psychological changes (Salthouse, 2011). Nevertheless, it is noteworthy that the MPFC is particularly vulnerable to aging functionally as the MPFC-DLPFC anticorrelation was statistically eliminated (also Wu et al., 2011).

3.2. Individual differences in WM capacity and MPFC-DLPFC

Greater MPFC-DLPFC anticorrelation at rest was associated with greater WM capacity across two independent samples totaling 97 young adults and using two different measures of WM capacity. In Experiment 1, greater WM capacity among young adults, as measured by a Letter-Number Sequencing task, was associated with greater right-lateralized MPFC-DLPFC anticorrelation. In Experiment 2, greater WM capacity among young adults, as measured by a composite score from the Operation-Span and Reading-Span tasks, was associated with greater left-lateralized MPFC-DLPFC anticorrelation. Importantly, all of these correlations were observed in an analysis in which MPFC-DLPFC anticorrelations were defined consistently and independently from the WM capacity measures. The observation that greater MPFC-DLPFC anticorrelation was associated with greater WM capacity regardless of the specific measure of WM capacity indicates that this brain-behavior relation is generalizable.

Among young adults, the laterality of the MPFC-DLPFC anticorrelation shifted across different measures of WM capacity, and it is unclear what factor explains this difference. All three WM capacity measures involved simultaneous maintenance and manipulation of information in WM, with variation in the amount of that information. An advantage of such complex tasks is that they tax WM capacity, but a disadvantage is all the tasks involve many kinds of mental operations. In Experiment 2, the primary WM measure was a composite of the Operation-Span and Reading-Span tasks. Such a composite has the virtue that it provides a task-independent latent measure of the underlying construct of WM capacity that is not overly sensitive to the measurement properties of a single task. It is also possible that different WM measures are related to different neural circuits. Overall, however, the two experiments converged in showing that greater magnitudes of MPFC-DLPFC resting-state anticorrelations were associated with greater WM capacity among young adults.

3.3. Methodological considerations

Compared to previous resting-state fMRI studies examining aging and individual differences, the present study had a number of potential methodological advantages. First, we...
used the aCompCor method of noise reduction (Behzadi et al., 2007) as implemented in Conn (Whitfield-Gabrieli & Nieto-Castanon, 2012). This method avoids explicit global signal regression, a widely used preprocessing technique known to mathematically generate anticorrelations (Murphey et al., 2009; Saad et al., 2012; Van Dijk et al., 2010; Wong, Olafsson, Tal, & Liu, 2012) which as a result renders anticorrelations uninterpretable (e.g., Chang & Glover, 2009) and may compromise the interpretability of positive correlations (Saad et al., 2012). The approach used in the present study is more likely to yield interpretable negative correlations and provides higher sensitivity and specificity for positive correlations (Chai et al., 2012). The differences in anticorrelations observed across age groups and individual differences within age groups observed in the present study are therefore less likely attributable to artifacts from data processing methods and may reflect biological processes. Second, we employed a method of artifact rejection above and beyond motion regression in order to reduce motion-related artifacts common in aging. As expected, the older adults had significantly more artifactual time points removed from analysis (although the percentage of time points removed was small for both groups). After artifact rejection, there was an apparent increase of posterior anticorrelations in the older adults, whereas frontal anti-correlations remained eliminated even at a liberal threshold of \( p = .05 \) uncorrected. These findings suggest that aging disproportionately degrades MPFC-DLPFC anticorrelations.

Greater movement in older relative to younger adults raised the possibility that differential connectivity findings could reflect differential movement. This appears unlikely, however, because age-related differences were also found when comparing younger and older groups after they were matched for movement and artifact outliers. Thus, the age-related differences are more likely to reflect actual differences in intrinsic brain organization.

Finally, the major findings regarding individual differences in intrinsic functional brain organization in relation to WM capacity were based on brain imaging correlations independent of behavioral measures. Specifically, the DLPFC regions were defined without reference to the WM behavior. Thus, the relations between the MPFC-DLPFC anticorrelations and WM performance were not contingent upon a whole-brain search for correlations with performance.

4. Limitations

One limitation to this study is that the older adult population may have substantially more cortical atrophy and more CSF, which may have differentially affected the normalization procedure. Another limitation is that although the use of different WM measures in Experiments 1 and 2 promote the generalizability of the findings, the different measures precluded a direct replication. Finally, anticorrelations are consistently found in young adults between specific neuro-anatomical systems (e.g., the DMN and the fronto-parietal control network) and the magnitude of these anticorrelations relate to individual differences in behavior (e.g., WM performance), but the neural mechanisms underlying these anticorrelations remain unknown.

5. Conclusion

In older adults, there was reduced WM capacity and the apparent elimination of MPFC-DLPFC anticorrelation. In younger adults, there were associations between greater magnitudes of WM capacity and greater MPFC-DLPFC anticorrelations. These results suggest that intrinsic anticorrelations between the MPFC, a node in the DMN, and DLPFC, a cortical region involved in cognitive control, may serve as a shared indicator of WM capacity both in aging and in individual differences among young adults. Also, just as WM capacity declines in older adulthood, WM capacity grows markedly in development from childhood to young adulthood, and so do MPFC-DLPFC anticorrelations (Chai, Ofen, Gabrieli, & Whitfield-Gabrieli, 2014).

Differences in intrinsic functional organization in the resting state may reflect the ongoing history of interactions among brain regions during active cognitive performance in everyday life. Ultimately, it will be valuable to relate directly such resting-state and active-performance network dynamics. Resting-state studies are limited in interpretation by the absence of ongoing behavioral measures. Conversely, active performance studies are limited by the interpretation of brain activations occurring at different levels of performance in younger and older adults. Thus, one study reported that age-related reductions in activation during a WM task can be understood essentially in terms of variation in WM capacity (Schneider-Garces et al., 2012). Another study reported that age-related differences activation cannot be explained solely by variation in WM capacity (Bennett, Rivera, & Rypma, 2013). Future studies that integrate resting-state and active-performance measures in younger and older adults with varying WM capacities in both age groups may clarify the extent to which individual differences among younger and older adults and age-related declines in older adults reflect shared and unique brain differences.

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