

Cerebral White Matter Integrity and Cognitive Aging: Contributions from Diffusion Tensor Imaging

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Abstract The integrity of cerebral white matter is critical for efficient cognitive functioning, but little is known regarding the role of white matter integrity in age-related differences in cognition. Diffusion tensor imaging (DTI) measures the directional displacement of molecular water and as a result can characterize the properties of white matter that combine to restrict diffusivity in a spatially coherent manner. This review considers DTI studies of aging and their implications for understanding adult age differences in cognitive performance. Decline in white matter integrity contributes to a disconnection among distributed neural systems, with a consistent effect on perceptual speed and executive functioning. The relation between white matter integrity and cognition varies across brain regions, with some evidence suggesting that age-related effects exhibit an anterior–posterior gradient. With continued improvements in spatial resolution and integration with functional brain imaging, DTI holds considerable promise, both for theories of cognitive aging and for translational application.

Keywords Diffusion tensor imaging · White matter · Cognition · Aging · Information processing · Human development

Research in areas defined traditionally as clinical neuropsychology, cognitive neuroscience, and cognitive aging has coalesced recently into a new discipline, the cognitive neuroscience of aging (Cabeza et al. 2005; Grady 2008). A major impetus in this research development has been the work conducted in structural and functional neuroimaging, especially positron emission tomography (PET), and magnetic resonance imaging (MRI). Imaging research has been extremely informative regarding the brain localization of cognitive function and age-related differences in brain structure and function. The majority of this work, however, has to date concentrated on cerebral gray matter. This is an understandable bias, given the importance of cortical activity for cognition. The remaining 40–50% of the brain volume, however, comprises white matter, regions where there is a preponderance of axons coated with myelin, which also contributes significantly to efficient cognitive functioning. Although some aspects of perceptual and cognitive performance are highly modular and localizable, others, such as attention and memory, depend on widely distributed neural systems. Thus, conditions leading to disconnection among these systems have significant consequences for behavior and cognitive functioning (Catani and Ffytche 2005; Filley 2005; Geschwind 1965a, b; Mesulam 1990).

Imaging studies of white matter volume have detected significant volumetric decline in healthy older adults, as compared to younger adults. Whereas the age-related decline in gray matter volume is relatively linear from younger adulthood, the corresponding decline in white matter

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tends to be nonlinear, with a plateau in middle-age and additional decline, beyond that of gray matter, in later adulthood (Guttman et al. 1998; Jernigan et al. 2001; Raz 2000; Raz et al. 2005; Salat et al. 1999).¹ Imaging studies of healthy adults also suggest an association between age-related decline in white matter volume and deficits in cognitive performance (Brickman et al. 2006), but the majority of previous research on the relation between white matter and cognition has been clinical studies of patient populations. In demyelinating diseases, such as multiple sclerosis (MS), many changes occur, from the level of visual psychophysical performance (Galvin et al. 1977; Regan et al. 1977), to information processing speed (Kail 1998; Litvan et al. 1988) and memory (Thornton and Raz 1997), although the effects of MS are not entirely uniform (Halligan et al. 1988; Jennekens-Schinkel et al. 1990; Rao 1995).

During normal aging, ischemic lesions, expressed as white matter hyperintensities (WMH) on MRI, are common among older adults and are correlated with decreased performance on some measures of cognitive performance (DeCarli et al. 1995; Gunning-Dixon and Raz 2000). These lesions occur in healthy adults, without significant impairment, although hypertension and cardiovascular disease appear to have some causal role (Raz et al. 2007). WMH tend to affect speed-dependent cognitive performance, and executive more than non-executive processing (Oosterman et al. 2004; Prins et al. 2005; van den Heuvel et al. 2006). There is some indication that WMH are more predictive of executive functioning decline when located in the frontal lobe (Gunning-Dixon and Raz 2003). From a series of regression analyses conducted on a sample of 65 individuals 65–84 years of age, Rabbitt et al. (2007) proposed that WMH prevalence accounted for all of the age-related variance in psychometric tests of speed and executive functioning. However, performance on more general measures of fluid intelligence (verbal and nonverbal problem solving, mental arithmetic) exhibited decline as a function of increasing age but no relation to WMH prevalence.

Diffusion Tensor Imaging

Research on cerebral white matter volume and WMH demonstrates that age-related differences in white matter

are relevant for understanding age-related effects in cognitive performance. A limitation of volumetric and lesion studies is that they provide little evidence regarding the microstructural properties of behaviorally relevant tissue. In addition, because cognition depends on the communication of widely distributed neural networks, information regarding the spatial organization and connectivity of white matter pathways would be valuable.

The development of diffusion tensor imaging (DTI) has provided a new avenue for understanding the relations among cerebral white matter integrity, cognitive functioning, and aging, and findings in this area have been accumulating rapidly (Johansen-Berg and Behrens 2009). DTI is a form of MRI that uses a tensor model to measure both the rate and directionality of the displacement distribution of water molecules across tissue components. Excellent discussions of the technical foundations and physics of DTI are presented in several sources: Bassler and Jones (2002), Beaulieu (2002), Jones (2008), Le Bihan (2003), Mori and Zhang (2006), and Mori (2007). For each voxel, DTI estimates diffusion in terms of the axes (eigenvectors) of an ellipsoid, typically separated into one major axis of diffusion and two minor axes that are orthogonal to the primary eigenvector. In gray matter, the diffusion of molecular water is highly isotropic—the same in all directions. In white matter, however, diffusion is more anisotropic—restricted to a particular direction. There are several sources of this directionality, including the myelin sheaths of axons, axonal cell membranes, and neurofilaments (Beaulieu 2002; Jones 2008; Peled 2007). Consequently, anisotropy is highest in white matter pathways with compact fiber bundles oriented in parallel, such as the corpus callosum and pyramidal tracts.

The most frequently used DTI summary measures are mean diffusivity and fractional anisotropy (FA). Mean diffusivity represents the average rate of diffusion, independent of the directionality, and is often expressed by the average of the apparent diffusion coefficients across the three orthogonal axes of the diffusion tensor. FA is a normalized (scalar) value that represents the fraction of the tensor that can be assigned to anisotropic diffusion. Because white matter with high structural integrity will typically place a greater degree of directional restriction on diffusion, increasing FA is frequently interpreted as reflecting a higher degree of white matter integrity. While this interpretation is a useful heuristic, it is important to recognize that the interpretation depends on the local architecture of white matter. When an imaging voxel contains crossing or kissing fibers, for example, then FA may be lowered even though axonal structural integrity is high (Pierpaoli and Bassler 1996; Virta et al. 1999).

Additional information regarding the neurobiological mechanisms of anisotropic diffusion can be derived from

¹ Virtually all neuroimaging studies of aging, as well as behavioral studies, are cross-sectional investigations of different age groups, rather than longitudinal studies of the same individuals over time. In describing cross-sectional results as *age-related*, we recognize that the differences among age groups include influences other than age (e.g., cohort effects). Thus, differences among the age groups in the dependent variables, expressed as deficits or improvements in particular measures, do not necessarily represent age-related *change*, for which longitudinal data are required.

the rate of diffusion (eigenvalue) along the individual eigenvectors. Axial diffusivity refers to the eigenvalue (λ_1) of the primary axis, whereas radial diffusivity is defined by the average of the eigenvalues perpendicular to the primary axis (λ_2, λ_3). Results from animal models suggest that myelin-specific damage tends to lead to increased radial diffusivity without significant changes in axial diffusivity, whereas axonal damage leads to decreased axial diffusivity (Song et al. 2002; Sun et al. 2006, 2007). This pattern is a signature of secondary (Wallerian) white matter degeneration following a primary lesion. The interpretation of the neurobiological mechanism of altered diffusivity, however, is not straightforward and, as with the interpretation of FA, depends on the local fiber architecture. In areas comprising pathways of intersecting fibers, rather than fibers arranged uniformly in parallel, secondary degeneration may lead to small or inconsistent changes in diffusivity. If, for example, within a region of intersecting pathways, such as the rostral pons, secondary degeneration primarily affects fibers of one direction (e.g., descending motor pathways), then the remaining fibers (e.g., transverse pontine fibers) would determine the apparent orientation of the principal eigenvector, possibly leading to a paradoxical increase in FA (Pierpaoli et al. 2001).

Age-Related Differences in DTI Measures of White Matter Integrity

Several reviews of DTI methods are available that specifically address the application to aging (Malloy et al. 2007; Moseley 2002; Sullivan and Pfefferbaum 2006, 2007; Wozniak and Lim 2006). We focus here on differences in the methods with which DTI data are extracted and analyzed, following image processing. These methods provide a context for the types of conclusions that are drawn, and each of the approaches has advantages and disadvantages for age-related comparisons.

Histogram analysis Histogram analysis provides an assessment of the frequency distribution of selected DTI measures across a range of voxels. In theory, the frequency distribution can be derived from any selected set of voxels (i.e., a region of interest, ROI), but histogram analysis has most often been applied to whole-brain data sets. Chen et al. (2001) proposed that whole-brain histogram analysis is both more accurate and less biased than an ROI-based approach. A single, whole-brain value is consequently attractive as a summary measure for comparison between groups. Several studies have reported an increase in mean diffusivity as a function of increasing adult age, derived from whole-brain histograms, suggesting an age-related decline in white matter integrity (Chen et al. 2001;

Nusbaum et al. 2001; Rovaris et al. 2003). Chen et al., however, found that the age-related increase was less pronounced for a tissue-specific estimate of mean diffusivity (1% per decade) than for a global estimate (3% per decade), implicating the potential contribution of cerebrospinal fluid (CSF) and large vessels to the global estimate. Whereas Z. G. Chen et al. considered anisotropy to be primarily a confounding influence in the measure of diffusivity, Rovaris et al. (2003) compared whole-brain FA and diffusivity histograms directly and found that although both types of histograms exhibited significant age-related effects, the age effect was strongest for FA, and the two histograms were not correlated. Rovaris et al. concluded that FA provides an independent index of white matter microstructure.

ROI analysis Information regarding regional, rather than whole-brain, differences in white matter integrity is valuable, especially when the goal is to correlate white matter measures with cognitive measures. Defining white matter ROIs in a reliable and accurate manner, however, is challenging. Due to the higher level of distortion in tensor images (from B_0 field inhomogeneity and eddy currents) than in T_1 -weighted anatomical imaging, ROIs drawn on T_1 -weighted anatomical images cannot be applied to tensor images without significant post-processing techniques. It is possible to define ROIs within the tensor images, but as Pfefferbaum et al. (2000) pointed out, that approach entails the danger of using the dependent variable (FA) to define the independent variable (anatomical ROIs).

Sullivan, Pfefferbaum, and colleagues developed a method of ROI analysis as a basis for their seminal series of DTI studies, which were the first to document programmatically the variation in white matter integrity associated with aging (Pfefferbaum and Sullivan 2003; Pfefferbaum et al. 2000; Sullivan et al. 2001). These authors used a tissue segmentation procedure to distinguish white matter, gray matter, and CSF in the tensor images, and then applied ROIs derived from anatomical imaging to the segmented DTI data. Pfefferbaum et al. (2000) assessed FA (among other measures) for five ROIs in a sample of 31 healthy men between 23 and 76 years of age. The correlation between FA and age was negative for all of the ROIs, except the splenium. Sullivan et al. (2001) compared these 31 men to 18 women between 23 and 79 years of age and found a similar pattern of age-related effects for men and women. Pfefferbaum and Sullivan (2003) applied systematic morphological expansion and restriction of selected brain regions and found that partial voluming effects had significant influences on DTI measures, but that the age-related differences (decline in FA, increase in diffusivity) could not be attributed entirely to partial voluming (see also (Bhagat and Beaulieu 2004)). Pfefferbaum et al. (2000)

concluded that the age-related decline in FA represented a decline in white matter integrity, possibly involving mild demyelination and loss of myelinated axons. These authors also suggested that the differential age-related effects in anterior regions were due to the later myelination of these regions.

Other methods of ROI analysis have yielded age-related differences consistent with those of Sullivan, Pfefferbaum, and colleagues (Abe et al. 2002; Nusbaum et al. 2001; O'Sullivan et al. 2001; Yoon et al. 2008). In several studies, Madden and colleagues have defined ROIs within the tensor images, in native space (Bucur et al. 2008; Madden et al. 2004, 2007). These authors reduced the potentially confounding effect of FA by lowering the maximal FA value in the image during ROI definition, so that relevant white matter structures were delineated as pure white. Data analyses are then performed with the FA limit removed. Gold et al. (2008) have also used this method. Figure 1 provides an example of ROIs defined in native space within the tensor images. The studies using this method generally confirm the previous findings, in that age-related decline in FA was typically greater in magnitude for anterior regions (e.g., genu, pericallosal frontal), although age-related decline was evident in posterior ROIs also.

Voxelwise analysis Defining ROIs anatomically has the advantage of using a common anatomical boundary for each participant, but the boundary definition depends on the judgment of individual operators. In addition, significant differences in the measures of interest may occur outside the selected ROIs. Alternatively, integration of DTI data with high-resolution anatomical images (or with the b_0 image from the DTI series) is possible, using normalization and smoothing methods similar to those used in functional imaging (Friston et al. 1995; Smith et al. 2004). Although

the transformation will entail the loss of some information, the DTI data for each participant are referenced to a standardized stereotaxic space. As a result, analyses can also be conducted on a voxelwise basis throughout the brain rather than limited to selected ROIs. Head et al. (2004), Salat et al. (2005), and Ardekani et al. (2007) conducted both voxelwise analyses and ROI analyses on spatially normalized DTI data. The results support the anterior–posterior gradient of age-related decline in white matter integrity (FA). Salat et al. also noted, however, that FA exhibited significant age-related decline in the posterior periventricular region. This latter region is a common location of WMHs in older adults (Holland et al. 2008; Vernooij et al. 2008).

A variant of the voxelwise approach, tract-based spatial statistics (TBSS) (Smith et al. 2006, 2007), provides more specific information about individual white matter tracts, without spatial smoothing. With the TBSS analysis, individual diffusion images are first aligned to a standard template. These images are then averaged across all participants to form a mean diffusion image that is used to generate a white matter “skeleton,” representing the center of white matter tracts that are shared by all participants (Fig. 2). The FA values within the skeleton can then be obtained for each participant. Several studies, using TBSS, have confirmed the anterior–posterior gradient of decline in FA that was observed with ROI and voxelwise methods, for healthy younger and older adults (Bennett et al. 2009; Burzynska et al. 2009; Damoiseaux et al. 2009).

DTI tractography Recently, considerable interest has been generated by DTI tractography (Catani 2006; Ciccarelli et al. 2008; Mori and van Zijl 2002; Nucifora et al. 2007; Wakana et al. 2004). Like voxelwise methods, a goal in tractography is to minimize the reliance on operator-defined parameters.

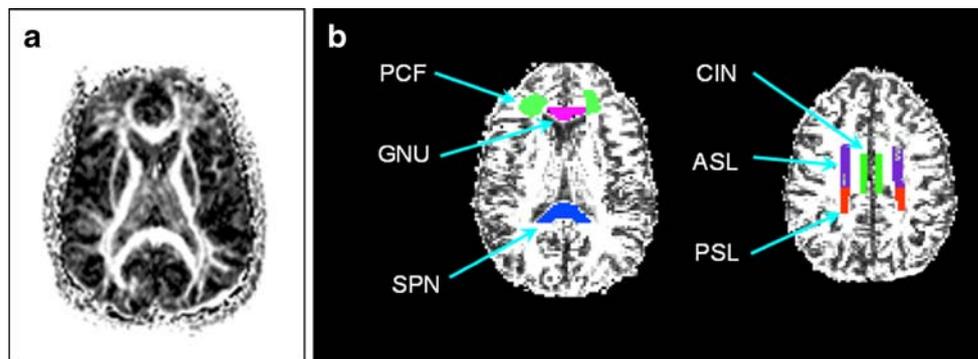


Fig. 1 Examples of regions of interest (ROIs) used in the analysis of the diffusion tensor imaging (DTI) data. **a** raw tensor image, with increasing fractional anisotropy (FA) represented as increasing brightness; **b** tensor images with maximum value for fractional anisotropy set to a lower threshold. Lowering the maximal FA threshold decreases the potential contribution of variation in FA to

the ROI definition. *PCF* pericallosal frontal; *GNU* genu of corpus callosum; *SPN* splenium of corpus callosum; *CIN* cingulum bundle; *ASL* anterior portion of the superior longitudinal fasciculus; *PSL* posterior portion of the superior longitudinal fasciculus. Figure modified from Chen et al. (2009) © *Brain Structure and Function* and Springer-Verlag Berlin Heidelberg, 2009

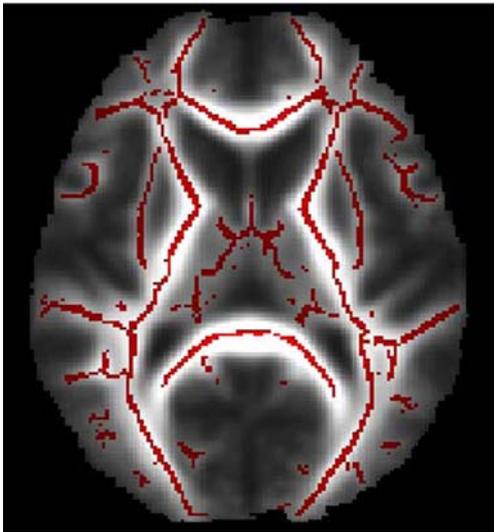


Fig. 2 Example of a white matter skeleton used in tract-based spatial statistic (TBSS) analyses. The white matter skeleton (red) represents the center of tracts common to all participants. It is superimposed on the mean diffusion image, which was created by averaging aligned FA diffusion images from each individual in the group. Figure modified from Bennett et al. (2009). © *Human Brain Mapping* and Wiley-Liss, Inc., 2009

But tractography places a higher premium on identifying the relevant white matter pathways within each participant, rather than on the degree to which the individual pathway is shared with the group average. Typically, tractography is an automated algorithm, conducted in native space, which estimates the most likely structure of an individual white matter pathway, either propagating from a specified source region or connecting source and target regions. Because tractography is conducted in native space, the spatial structure of the tracts will vary across participants.

Tractography algorithms can be broadly categorized as being either deterministic or probabilistic (Jones 2008; Nucifora et al. 2007). Deterministic algorithms reconstruct tracts by estimating a trajectory from each seed point along a streamline that is estimated from the primary diffusion directions of adjacent voxels (Basser et al. 2000; Conturo et al. 1999; Mori et al. 1999). In quantitative tractography (Corouge et al. 2006; Sullivan et al. 2006, 2008), FA and diffusivity values are obtained continuously (e.g., at millimeter intervals) along the length of the tract. With probabilistic algorithms, in contrast, trajectories follow a probabilistic distribution of potential primary diffusion directions that are estimated for each voxel (Behrens et al. 2003; Parker et al. 2003). Probabilistic tractography is valuable for tracking into regions with low anisotropy, such as cortical and subcortical gray matter and areas with crossed or kissing fibers (Jones 2008; Nucifora et al. 2007).

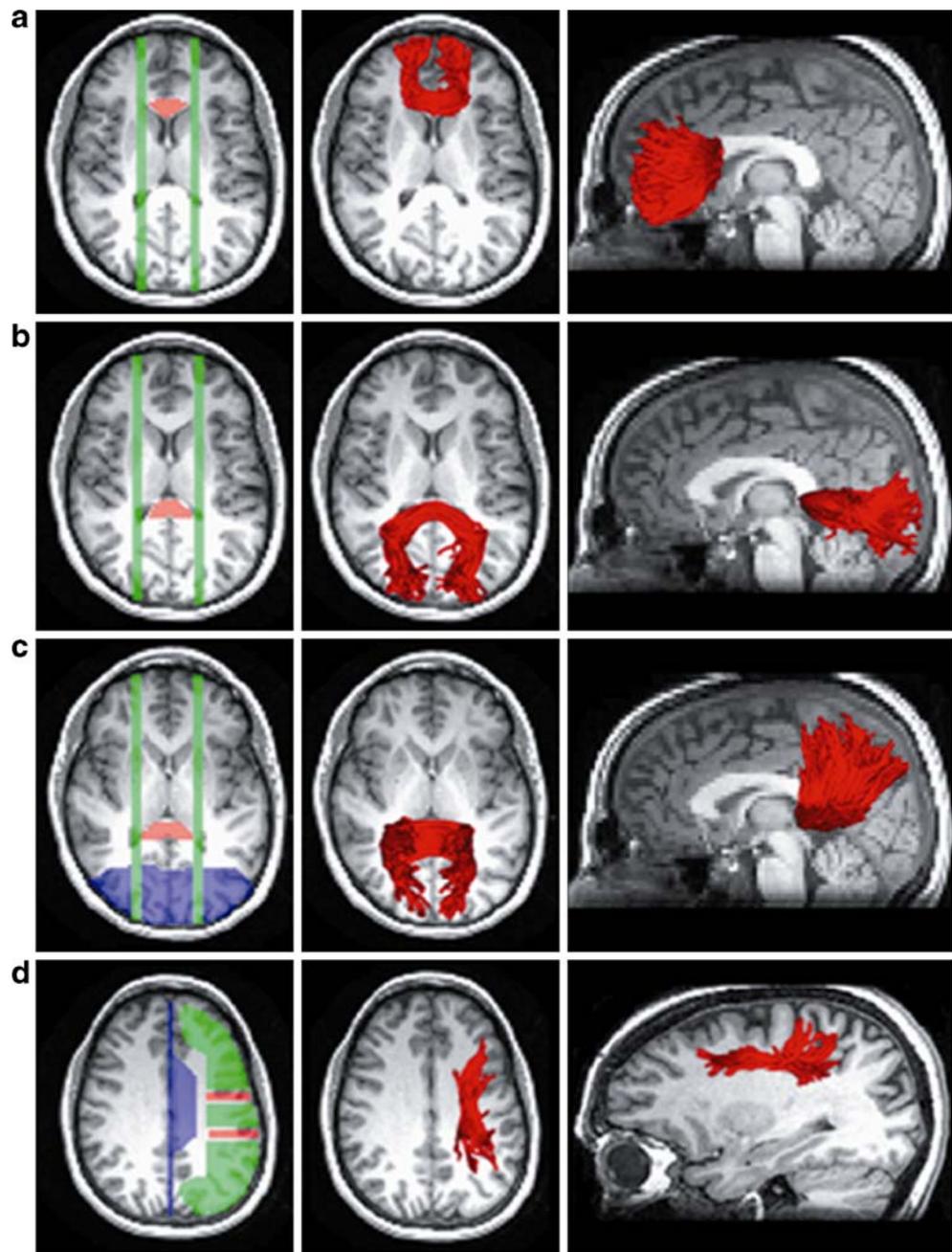
Overall, DTI tractography has been successful in isolating white matter pathways with a remarkably high

degree of correspondence to classical, post-mortem dissection and histology (Catani et al. 2002; Conturo et al. 1999; Dauguet et al. 2007; Lawes et al. 2008). To date, investigations of adult age differences using tractography have relied primarily on deterministic methods. Sullivan et al. (2006) reported the first application of fiber tracking to aging. These authors focused on the corpus callosum, using the high-resolution T_1 -weighted images to create an averaged image space for 10 younger and 10 older adults combined. Thus, target and source regions could be defined within a common anatomical space and transformed back into native space for quantitative fiber tracking. Fibers were estimated across six sections spanning the anterior–posterior length of the corpus callosum, yielding within-section estimates of number of fibers, fiber length, FA and mean diffusivity. The two most anterior sections of the callosum exhibited significant age-related decline in FA and increase in diffusivity, whereas age differences were more variable for the measures of fiber number and length.

Subsequent applications of quantitative fiber tracking to other white matter pathways have generally confirmed the age-related pattern reported by Sullivan et al. (2006) for callosal fibers (Davis et al. 2009; Madden et al. 2009; Sullivan et al. 2008; Zahr et al. 2009). Figure 3 presents an example of deterministic tractography, from Madden et al. (2009). The FA data averaged across 20 younger and 20 older adults are presented along the length of four separate tracts in Fig. 4. These data, with healthy, community-dwelling individuals, reveal that age-related decline in FA is most apparent in frontal pericallosal regions, consistent with the anterior–posterior gradient of previous studies. Sullivan et al. (2008), however, completed tractography on 11 bilateral pathways, in addition to the callosum, and proposed that a superior–inferior gradient is also evident. Greater age-related decline occurred in the microstructure of frontal, limbic, striatal, and superior pathways of the supratentorium, as compared to infratentorial (pontine and cerebellar) systems. Zahr et al. (2009), in a fiber tracking study of eight pathways, have also confirmed that both anterior–posterior and superior–inferior gradients were present.

Axial and radial diffusivity As noted previously, the commonly used DTI summary metrics FA and mean diffusivity are calculated from the individual eigenvalues of the tensor matrix, λ_1 , λ_2 , λ_3 , which in turn comprise axial diffusivity (λ_1) and radial diffusivity (mean of λ_2 and λ_3). Examining the individual eigenvalues can provide information regarding the neurobiological mechanisms of age-related effects in the summary metrics. There is some evidence, for example, that the age-related differences in FA and mean diffusivity are driven primarily by an increase in radial diffusivity (Bhagat and Beaulieu 2004; Madden et

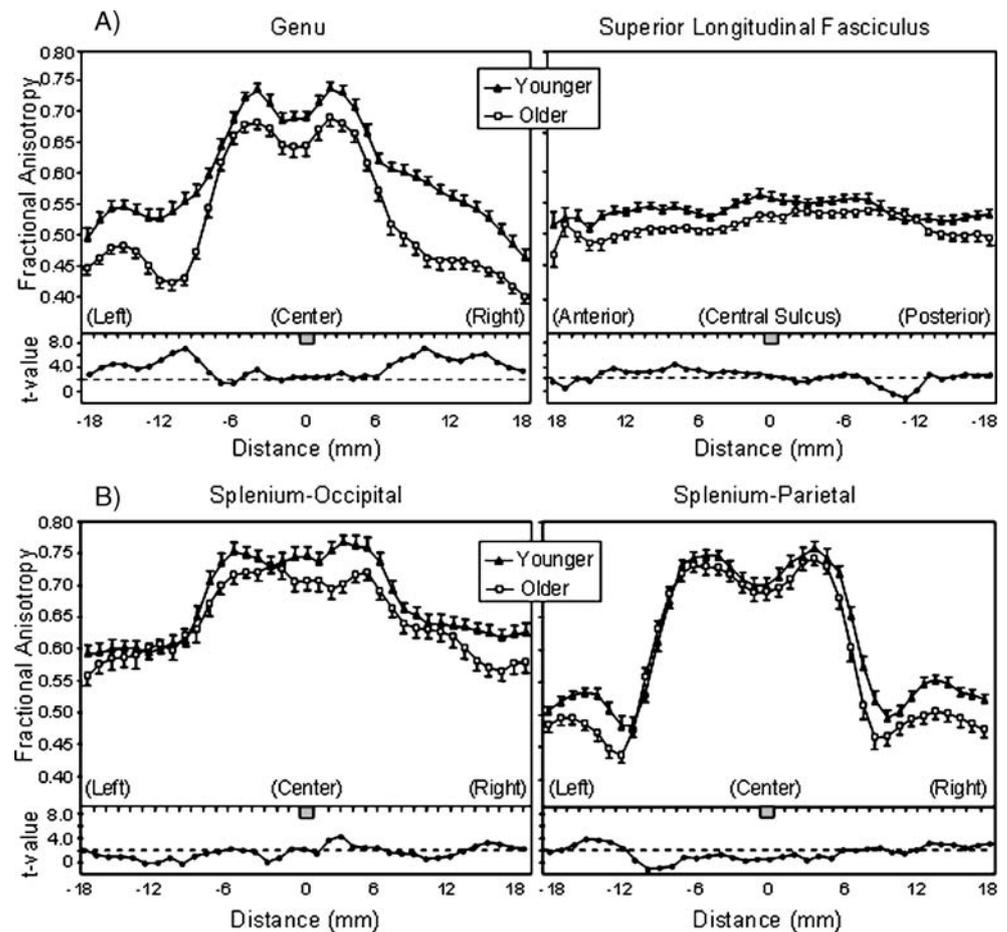
Fig. 3 Examples of deterministic tractography. Fiber tracts (in red) generated by target and source region placement, for a single participant. The orange areas are target regions, the green areas are source regions, and the blue areas are exclusion regions. All of these regions are operator defined, for each participant, using anatomical boundaries. The fiber tracking algorithm estimates tracts that pass through the target regions from the source regions, eliminating any fibers terminating in the exclusion regions. The approximate locations of output fiber tracts are illustrated by overlaying on a single-slice T₁-weighted image. **a** genu; **b** splenium-occipital; **c** splenium-parietal; **d** superior longitudinal fasciculus. Figure modified from Madden et al. (2009). © *Journal of Cognitive Neuroscience* and MIT Press, 2009



al. 2009; Zhang et al. 2008). Given the association of radial diffusivity with myelin-related effects (Song et al. 2002; Sun et al. 2006, 2007), an age-related increase in radial diffusivity suggests a differential contribution of demyelination (i.e., a thinning or deterioration of the axonal myelin sheath) to age-related decline in FA. Analyses focusing on the regional changes in axial and radial diffusivity suggest that the differential age-related increase in radial diffusivity is more pronounced in the prefrontal regions. That is, the anterior–posterior gradient of FA may reflect a comparable gradient of declining myelin integrity (Davis et al. 2009; Sullivan et al. 2006; Zahr et al. 2009).

Other patterns of age-related differences in axial and radial diffusivity, however, have been observed. Sullivan et al. (2008), Vernooij et al. (2008), and Zahr et al. (2009) reported significant age-related increases in both the axial and radial components of diffusivity, across a wide selection of white matter pathways. Using TBSS, Bennett et al. (2009) and Burzynska et al. (2009) categorized various white matter regions according to the relative change in the axial and radial components of diffusivity. In both of these latter studies, radial diffusivity increased with age across the majority of tracts, but the occurrence of a concomitant increase in axial diffusivity was more

Fig. 4 Mean FA as a function of age group and interval along the tract. *Error bars* represent 1 *SE*. **a** genu and superior longitudinal fasciculus; **b** splenium-occipital and splenium-parietal. For genu and splenium, the tracts are oriented *left–right*, with 0 = axial midline. For the superior longitudinal fasciculus, the tracts are oriented anterior–posterior and 0 = central sulcus. Below the mean FA data, *t*-values are plotted for the age group comparison at each point along the tract. The *dotted line* represents the significant *t* value for $p < .05$, two-tailed. Figure modified from Madden et al. (2009). © *Journal of Cognitive Neuroscience* and MIT Press, 2009



variable. The genu of the corpus callosum consistently exhibited age-related increases in both axial and radial diffusivity, whereas increased diffusivity was more likely to be limited to the radial component in association tracts connecting anterior and posterior regions (e.g., superior longitudinal fasciculus). Further, both Bennett et al. and Burzynska et al. found that, in some white matter tracts, aging was associated with decreased axial diffusivity.

These results suggest that an age-related decrease in myelin integrity alone cannot account for the pattern of age-related differences in axial and radial diffusivity. The relevant neurobiological mechanisms are not currently known. Pre-frontal regions are among the last to myelinate during development, and thus a “last-in-first-out” principle may contribute to the observed anterior–posterior gradient (Bartzokis et al. 2004; Davis et al. 2009; Pfefferbaum et al. 2000; Raz 2000). Myelin is only one of the variables, however, leading to restricted diffusion (Beaulieu 2002; Jones 2008). Decreased density of axonal packing within a voxel, due to decreased axonal diameter, fewer axons, or both, would also lead to increased diffusivity in all directions (Sullivan et al. 2008; Zahr et al. 2009). Decreased axial diffusivity may represent an initial response to lost axons

(i.e., increased extracellular matrix and glia), which is followed by increased axial diffusivity as cellular debris is cleared by microglia (Burzynska et al. 2009). Decreased axial diffusivity may also reflect macrostructural effects, such as a decreased coherence in the orientation of axons, within voxels, for older adults relative to younger adults (Bennett et al. 2009). That is, for two axonal bundles of comparable structural integrity, the bundle in which the axons are less consistently aligned will have lower axial diffusivity.

A related issue, emphasized by Wheeler-Kingshott and Cercignani (2009), is that an individual eigenvalue does not necessarily reflect the same biophysical substrate across data sets, if the data sets differ in the orientation of the principal eigenvector. Whether changes in the individual eigenvalues, which are properties of the tensor, reflect true changes in the tissue microstructure, depends on the mathematical and geometrical properties of the data, especially the orientation of the principal eigenvector. Thus, axial and radial diffusivity are important as additional sources of information regarding differences across DTI data sets, but interpreting the neurobiological substrate of these variables is limited, without determining the orientation of the principal eigenvector.

Cognitive Correlates of White Matter Integrity in Aging

Identifying the cognitive correlates of age-related differences in white matter integrity has been a central concern of DTI research (Gunning-Dixon et al. 2009; Moseley 2002; Sullivan and Pfefferbaum 2006, 2007). This goal, however, depends on the available data regarding the relation between white matter integrity and cognition in healthy adults, independently of age effects. The relation between white matter and cognition has not yet been established clearly, and though consistent trends are evident, no single classification scheme adequately characterizes the extant findings.

Speed of Processing and Executive Functioning As discussed previously, some indication of a relation between white matter integrity and cognition derives from research on ischemic lesions of white matter, which have demonstrated that increasing WMH volume is associated with decreasing performance on tests of elementary perceptual speed (Rabbitt et al. 2007; Raz et al. 2007; van den Heuvel et al. 2006). One general principle that emerges from DTI studies is that, independently of age effects, decreasing integrity of normal appearing white matter is associated with worse performance on tests that rely on processing speed and executive functioning. This relation has been confirmed with both voxelwise (Turken et al. 2008), and tractography-based (Correia et al. 2008) DTI techniques. The relation between white matter integrity and speed also extends to tasks such as lexical decision (word/nonword discrimination), which rely on the retrieval of semantic memory information (Gold et al. 2007).

Although most measures of processing speed are defined on the basis of manual reaction times and thus have a significant motor component, the white matter-speed relation appears to hold for information processing stages that precede the motor response. Gold et al. (2007) found that, in younger adults, decreased white matter integrity (FA) in language regions of the left hemisphere (inferior frontal and inferior parietal) was correlated with slower lexical decisions, but that a corresponding effect was not evident in regions mediating visual/motor processing (optic radiation and posterior limb of the internal capsule, bilaterally). Similarly, data from combined imaging modalities of DTI and magnetoencephalography (MEG) suggest that the correlation between white matter integrity and the latency of peak visual responses during an eye movement task, in younger adults, holds at an early stage of information processing, within the first 120 ms of saccade initiation (Stufflebeam et al. 2008).

Vernooij et al. (2009) provided a particularly strong source of support for the relation between white matter integrity and processing speed in a population-based

sample of 860 older adults 61–92 years of age (Rotterdam Study). These authors conducted regression analyses of whole-brain DTI variables and cognitive performance that statistically controlled the relative volume of both normal appearing white matter (i.e., atrophy effects) and white matter lesions. A variety of cognitive measures yielded composite measures of information processing speed, motor speed, memory, and executive functioning (e.g., response fluency and inhibition). Declining FA of normal appearing white matter was related significantly to declines in both motor speed and processing speed, but only the latter relation held for mean FA when lesion volume was controlled. In addition, both axial and radial diffusivity exhibited more widespread relation, than mean FA, to the cognitive measures. Both diffusivity measures were correlated significantly with information processing speed and executive functioning, with lesion volume controlled (i.e., higher diffusivity associated with worse performance). None of the DTI variables was related to memory performance. Vernooij et al. also used age as a covariate in their analyses, however, and thus the observed correlations between white matter integrity and processing speed were independent of age-related variability in processing speed.

Aging, White Matter Integrity, and Cognition: Interpretive Issues The DTI studies reviewed to this point lead to two conclusions. First, age-related differences occur both in the overall measures of white matter integrity (e.g., FA) and in component measures (e.g., axial and radial diffusivity), especially in prefrontal regions. Second, independently of age, variation in white matter integrity is correlated with cognitive performance, particularly in tests relying on speed of information processing and executive functioning. An independent question is whether there are interactive effects in the aging-white matter and cognition-white matter trends. Although individual research studies bearing on this question have been accumulating rapidly, several issues should be considered in interpreting the results.

The most fundamental question to be addressed is the causal role of white matter integrity in age-related differences in cognitive performance. Specifically, to what extent is age-related variability in cognitive performance shared with age-related differences in white matter integrity? To date, however, few studies have conceptually framed the research in this manner. Individual studies have addressed a variety of different, though related, issues that do not always map directly onto this fundamental question. For example, the demonstration of a correlation between white matter integrity and some cognitive measure, for older adults, does not necessarily imply that white matter integrity is associated with the *age-related* variance in the cognitive measure. Similarly, if age groups are analyzed

separately, and a correlation between white matter integrity and cognition is statistically significant for older adults but not for younger adults, this pattern, by itself, does not necessarily imply that the correlation differs significantly between the age groups. The correlation may be slightly above threshold for older adults, but slightly below threshold for younger adults. A direct comparison of the age groups, or some test of the age-related difference in the relation between the DTI and cognitive measures, is necessary.

In addition, although both the cognitive and DTI measures may vary with age, this pattern is not sufficient to infer that white matter integrity has a direct influence on age-related effects in cognitive performance. To infer a mediating or causal role, the DTI measures should have a significant relation to the measures of cognition that is independent of age, and the age-related variance in the cognitive measure should be attenuated by including the DTI measures in the regression model predicting cognition from age (Baron and Kenny 1986; Lindenberger and Pötter 1998; Salthouse 1992).

Related to these methodological concerns is the issue of identifying specific cognitive effects in aging. Although the general trend of DTI studies suggests that the pronounced influence of white matter integrity is related to processing speed and executive functioning (Correia et al. 2008; Stufflebeam et al. 2008; Turken et al. 2008; Vernooij et al. 2009), it is not clear whether age-related differences in these aspects of cognitive functioning are entirely separable at the behavioral level. Some behavioral studies suggest that age-related differences in executive functioning are empirically separable from speed-dependent differences (Keys and White 2000; Rodriguez-Aranda and Sundet 2006), and the anterior–posterior gradient that has been noted in DTI studies of aging and white matter integrity (Davis et al. 2009; Madden et al. 2009; Sullivan and Pfefferbaum 2006, 2007) is consistent with the assumption of an identifiable role of executive functioning. Elementary measures of information processing speed, however, share age-related variance with a wide variety of cognitive tasks (Madden 2001; Salthouse 1996; Salthouse and Madden 2007), including executive functioning (Salthouse et al. 2003), and neuropsychological tests that are assumed to measure frontal lobe functioning correlate highly with non-frontal tests (Salthouse et al. 1996). Thus, interpreting age-related variation in white matter integrity in terms of the anterior–posterior gradient and executive functioning may be a useful starting point for DTI research but ultimately is not likely to account for all of the findings (Bennett et al. 2009; Greenwood 2000; Kennedy and Raz 2009; Madden et al. 2009).

Finally, the potential role of health-related variables, in the interaction of age-related change in the brain and

cognition, is not well understood. As noted previously, the relation of DTI measures to cognitive performance is influenced by the presence of WMH (Vernooij et al. 2009), and WMH tend to increase as a function of both age and cardiovascular risk (e.g., hypertension). Thus, variation in DTI measures of white matter integrity, across age groups, is likely influenced by WMH and cardiovascular health.

Nitkunan et al. (2008), for example, examined the relation between white matter integrity measures from DTI and brain biochemistry from magnetic resonance spectroscopy (MRS), within a single ROI (white matter of the centrum semiovale). These authors examined cerebrovascular status by comparing three groups of older adults: individuals with cerebral small vessel disease (SVD), hypertensive individuals without a history of stroke, and normotensive controls. Within each group, the DTI measures correlated significantly with MRS measures indicating axonal loss/dysfunction, and the magnitude of the correlations was generally graded with the severity of cerebrovascular disease, with the hypertensive group intermediate between the normotensive and SVD groups. The Nitkunan et al. analyses, however, did not focus specifically on the age-related variation in white matter integrity; age-related effects were covaried. Vernooij et al. (2008), examined the age-related effects and found that, in their Rotterdam Study participants, few age-related differences in FA remained significant once white matter atrophy and WMH lesion load were controlled statistically. Vernooij et al. concluded that age-related differences in white matter integrity reflect a pathophysiologic process rather than aging per se.

Alternatively, cardiovascular status and WMH volume may alter the normal relation between adult age and white matter integrity rather than entirely subsuming the relation between age and white matter integrity. Correia et al. (2008) derived several quantitative metrics of white matter integrity, based on deterministic tractography, and compared individuals with known vascular white matter injury (40–79 years) to a control group of demographically similar, healthy individuals (44–84 years). Metrics calculated from whole-brain data indicated significantly lower white matter integrity for the vascular group than for the controls. In addition, the metrics exhibited significant age-related decline for the healthy group but not for the vascular group, suggesting that subcortical vascular disease disrupted the normal variation in white matter integrity with age.

Aging, White Matter Integrity, and Cognition: Empirical Findings The results of DTI studies, regarding cognitive correlates of age-related differences in white matter integrity, have yielded a complex pattern. But the data generally incorporate the trends that we have noted for 1)

the relation between aging and white matter (anterior–posterior gradient); and 2) the relation between cognition and white matter (correlation with speed and executive functioning).

O’Sullivan et al. (2001) reported one of the initial correlations between DTI and cognitive data for older adults. These authors used relatively broad ROIs (whole-brain white matter divided into anterior, middle, and posterior regions) and correlated mean diffusivity and FA with several psychometric tests of cognitive function. Correlations performed within the older adult group indicated that mean diffusivity in the anterior ROI was correlated with a neuropsychological measure of executive functioning (Trail Making), whereas FA in the middle region was correlated with verbal fluency. These analyses, however, were conducted only within the older group, and the effect of age was covaried. O’Sullivan et al. suggested that the age-related decline in white matter integrity would lead to a disconnection of neural networks necessary for efficient cognitive performance. This concept of disconnection has been discussed widely as an explanatory construct for age-related cognitive decline, although whether the relevant neurobiological mechanism is demyelination, axonal loss, macrostructural organization, or a combination of these, is being investigated currently (Andrews-Hanna et al. 2007; Bartzokis 2004; Bartzokis et al. 2004; Charlton et al. 2006, 2008; Damoiseaux, et al. 2009).

Contemporaneously with the O’Sullivan et al. (2001) report, Sullivan et al. (2001) confirmed that white matter integrity was correlated with a behavioral measure, for 51 participants across a wide range of 23–79 years. A sensory/motor task requiring interhemispheric transfer (alternating finger tapping) correlated positively with mean FA, whereas a baseline version of the task (unimanual finger tapping) did not. Two aspects of the Sullivan et al. findings are particularly relevant: First, the positive correlation between FA and alternating finger tapping held within posterior ROIs (splenium and parietal pericallosal), whereas the age-related decline in mean FA was greater for more anterior regions. Second, in a regression model containing both age and FA, from the posterior ROIs, predicting finger tapping output, FA remained a significant predictor whereas age did not. This finding suggests that white matter integrity may have a more direct influence than age on this form of sensory/motor performance.

Madden et al. (2004) reported a similar pattern in an ROI analysis of 16 younger and 16 older adults. Mean reaction time in a choice response task (visual oddball) correlated with splenium FA for younger adults, but with FA within the anterior limb of the internal capsule for older adults. Rather than comparing the effectiveness of age and FA as predictors, Madden et al. took the approach of testing the age group difference in the correlation between mean

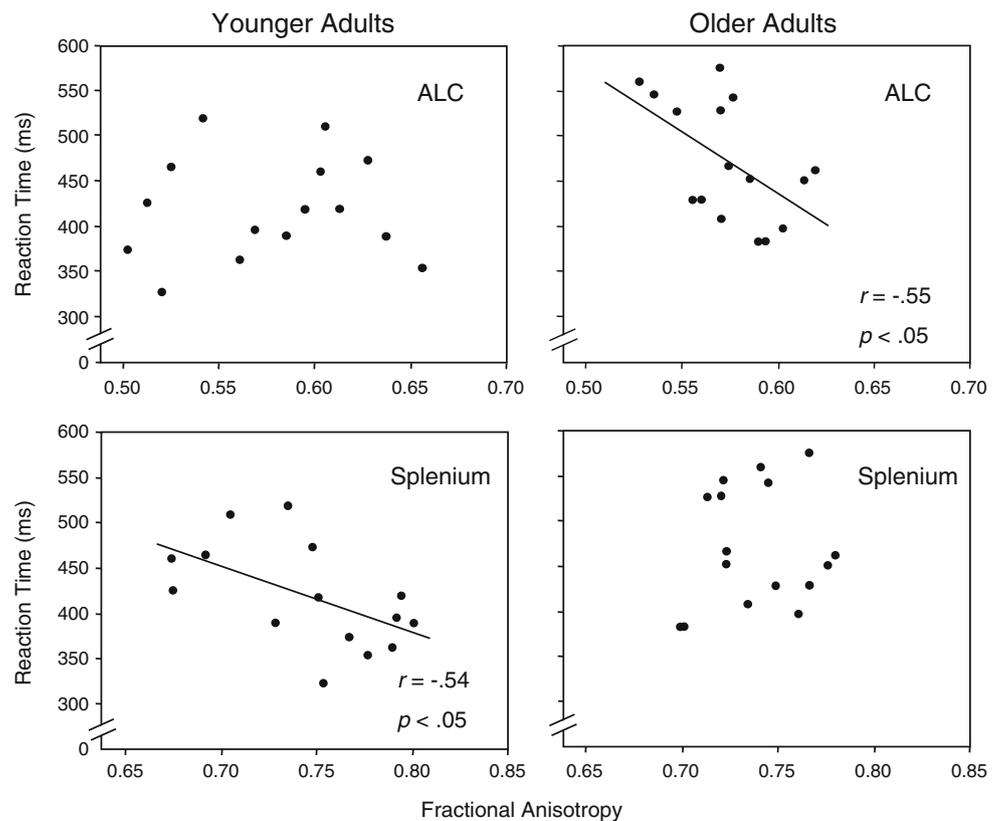
reaction time and FA. This comparison, illustrated in Fig. 5, was significant, which provided the first direct test of an age-related difference in the correlation between white matter integrity and a behavioral measure of cognitive performance. In addition, as in the Sullivan et al. (2001) study, those regions exhibiting significant correlations with behavioral measures were not those exhibiting the most pronounced age-related decline in mean FA.

Several subsequent studies have also focused specifically on age-related differences in the relation between white matter integrity and cognition. In their tractography study of younger and older adults, Davis et al. (2009) used a regression model that included the interaction between age group and FA as a predictor of each of several neuropsychological tests. These authors found that several tracts exhibited a stronger correlation between FA and cognitive performance for older adults than for younger adults. Specifically, older adults’ increasing FA within frontal regions (genu, uncinate fasciculus) was correlated with better performance on several tests of executive functioning (e.g., spatial working memory, set shifting), whereas the corresponding effect within older adults’ more posterior brain regions (e.g., splenium, inferior longitudinal fasciculus) reflected increasing FA and better performance on visual memory tests. Further, when the components of FA were distinguished, the correlations were nearly entirely due to radial diffusivity rather than axial diffusivity.

Similarly, Kennedy and Raz (2009), in their ROI analysis, also used regression models that tested for age-related differences in the relation between FA and several composite measures of cognitive performance. The regional variation in the correlations, however, was somewhat different than that reported by Davis et al. (2009). In the Kennedy and Raz data, age-related decline in white matter integrity within more anterior brain regions was associated with decreased processing speed and working memory, whereas decline within more posterior brain regions was associated with reduced inhibition and task switching. Kennedy et al. emphasized that the targeted cognitive domains, though differentiated into speed, executive functioning, and several forms of memory, still relied on widely distributed white matter pathways, in which disruption can have wide-reaching effects.

In recent studies, researchers have taken the further step of identifying the age-related variance in cognition and estimating the degree to which age-related variance in DTI is shared with age-related variance in cognition. Madden et al. (2009) and Gold et al. (2008) were both concerned with specific components of information processing. Madden et al. used a model of reaction time distributions to distinguish the efficiency of retrieval of semantic information (drift rate), in a word categorization task, from the more

Fig. 5 Relation between choice reaction time and FA, for younger and older adults. ALC = anterior limb of internal capsule. Figure modified from Madden et al. (2004). © *Neuroimage* and Elsevier, 2004



peripheral processes of display encoding and response time. In a series of regression analyses, age-related variance in the drift rate measure was attenuated substantially by individual differences in FA within regions of frontoparietal white matter pathways (central genu and splenium-parietal). Using a similar application of regression models, Gold et al. focused on the time required to switch between different tasks (letter or number decisions), rather than on the efficiency of individual decisions. Gold et al. found that FA within a frontoparietal pathway, the superior longitudinal fasciculus of the left hemisphere, was a significant mediator of age-related variance in task switching.

Zahr et al. (2009) examined domains of cognition: working memory, motor performance, and problem solving, rather than specific information processing components. As in the Madden et al. (2009) and Gold et al. (2008) studies, Zahr et al. found that the DTI measures led to significant attenuation of age-related variance in the cognitive measures. The genu and fornix were mediators of all three cognitive domains, whereas age-related differences in motor performance were influenced by a wider range of pathways, including the splenium and uncinate fasciculus in addition to the genu and fornix.

Charlton et al. (2008) used structural equation modeling to determine whether mean diffusivity within a relatively large ROI (primarily centrum semiovale) was a mediator of the relation between age and several measures of cognitive

performance: speed, cognitive flexibility, working memory, and fluid intelligence, for 118 adults 50–90 years of age. Surprisingly, although speed mediated the age-related effects in all of the other cognitive measures, diffusivity was a significant mediator only of the age-related variance in working memory. The authors obtained an identical model using mean FA rather than diffusivity as the DTI mediator.

As a preliminary, semi-quantitative roadmap through the diverse findings of DTI studies on aging and cognition, we have listed in Table 1 the results of 18 published reports of a statistical relation between one or more DTI variables and a behavioral measure of cognition, with a sample of older adults. We selected these studies from PubMed searches using the terms “white matter integrity AND aging” and “diffusion tensor imaging AND aging AND cognitive.” The selected articles used DTI to assess white matter integrity, examined white matter-cognition effects in healthy older adults, were not review papers, and used regional rather than whole-brain DTI measures. The table values are the mean effect sizes (Rosenthal and DiMatteo 2001) for the white matter-cognition relation within a brain region. We have used bold font for effect sizes that are moderate or larger ($> .30$). Note, however, that these effect sizes refer to the overall effect of the white matter-cognition relation, in studies of older adults, not to the age-related variation in this effect. In addition, the values are likely underestimates

Table 1 Summary of aging studies examining relationships between fractional anisotropy (FA) and cognitive functioning

	Frontal			Middle					Superior			Posterior		
	Front	aPERI	CCg	Mid	ALIC	PLIC	UNC	Temp	CING	CS	Par	CCs	pPERI	Occ
Executive functioning														
Composite measures (g, n, r)	0.19	–	0.80	0.05	–	–	0.35	0.34	0.29	–	–0.26	0.16	–	–
Working memory (h, j, o, p)	0.01	0.09	0.11	–	0.15	0.0	0.57	0.11	0.0	0.41	–0.08	0.0	0.28	–0.01
Set shift/task switch (a, j, l, o, p)	0.01	0.16	0.0	0.09	0.0	0.0	0.24	–0.12	0.0	–	0.05	0.0	–	–0.03
Memory span (j, o, p)	0.17	–	0.20	–	0.0	0.0	0.0	0.0	0.0	–	0.01	0.0	–	0.02
Fluency (a, c, f, h)	–0.05	0.17	–	0.69	–	–	–	–	–	0.20	0.37	–	0.16	0.22
Inhibition (p)	0.0	–	0.0	–	0.0	0.0	–	0.0	–	–	0.21	0.0	–	0.0
Processing speed														
Composite measures (g, p, r)	0.18	–	0.13	0.09	0.0	0.0	0.65	0.55	0.43	–	–0.06	0.06	–	0.0
Digit symbol (h, m)	0.30	0.14	0.14	0.10	0.14	–	–	–0.01	0.14	0.36	0.23	0.26	0.10	0.25
Simple/choice RT (d, h, j)	–0.13	0.15	0.0	–	0.31	–	–	–0.05	–	0.43	–0.19	0.0	0.01	–0.04
Finger tapping (b, j)	0.18	0.0	0.0	–	–	–	–	0.28	–	0.0	0.17	0.56	0.61	0.18
Interhemispheric processing (e)	–	–	0.48	–	–	–	–	–	–	–	–	0.51	–	–
Finger movements (m)	0.19	–	0.05	0.18	0.35	–	–	0.03	0.06	–	0.24	0.05	–	0.21
Memory														
Recognition (k, o)	–	0.49	0.16	–	–	–	0.08	0.0	0.22	–	–0.03	0.35	–	–
Free recall (f, n, p)	0.00	–	0.0	–	0.23	0.10	–	0.41	–	0.12	0.0	0.0	–	0.0
Paired-associate learning (o, p)	0.0	–	0.0	–	0.0	0.27	0.0	0.20	0.47	–	0.0	0.0	–	0.0
Mental ability														
Reasoning (c, f, h)	0.10	0.28	–	–	–	–	–	–	–	0.16	–	–	0.23	0.10
Irregular word reading ability (c, f, h)	0.04	0.08	–	–	–	–	–	–	–	0.23	–	–	0.03	0.07
Global cognitive functioning (c, f, h)	0.03	0.25	–	–	–	–	–	–	–	0.30	–	–	0.22	0.06
Word reading (i, q)	–	–	0.55	–	–	–	–	–	–	–	0.0	0.38	–	–
Postural stability														
Balance (b, m)	0.32	0.0	0.18	0.03	0.15	–	–	–0.03	–0.06	0.63	0.13	0.23	0.77	0.04

Values are mean effect sizes (Pearson r) for effects reported in the following studies: (a) O'Sullivan et al. (2001), (b) Sullivan et al. (2001), (c) Shenkin et al. (2003), (d) Madden et al. (2004), (e) Schulte et al. (2005), (f) Shenkin et al. (2005), (g) Charlton et al. (2006), (h) Deary et al. (2006), (i) Sullivan et al. (2006), (j) Grieve et al. (2007), (k) Bucur et al. (2008), (l) Gold et al. (2008), (m) Sullivan et al. (2008), (n) Ziegler et al. (2008), (o) Davis et al. (2009), (p) Kennedy and Raz (2009) (q) Madden et al. (2009), and (r) Zahr et al. (2009). Moderate effect sizes and larger (> 0.30) are listed in bold font

Mean effect sizes were computed following the method of (Rosenthal and DiMatteo 2001). Pearson r values were obtained from F and Spearman ρ statistics (in the latter case via conversion to standard normal deviate Z), averaging r values after transforming them to weighted Fisher z scores, and giving an r value of zero to effects described as not statistically significant. In some cases, r values were adjusted such that positive relationships indicate where better performance is associated with higher FA, and age-controlled r values were used when available. Data from Charlton et al. (2008), Thomas et al. (2008), and part of Sullivan et al. (2001) were not included because there was not enough information to obtain Pearson r values. RT reaction time

White matter tracts and regions were grouped into four broadly defined areas (Frontal, Middle, Superior, Posterior). *Front* frontal white matter (including forceps minor); *aPERI* anterior pericallosal white matter; *CCg* genu of the corpus callosum (including prefrontal, premotor/precentral, and postcentral subregions of the corpus callosum); *Mid* middle white matter (including fornix and external capsule); *ALIC* anterior limb of the internal capsule; *PLIC* posterior limb of the internal capsule; *UNC* uncinate fasciculus; *Temp* temporal white matter (including inferior longitudinal fasciculus); *CING* cingulum bundle; *CS* centrum semiovale (including large regions of interest from slices of superior white matter); *Par* parietal white matter (including superior longitudinal fasciculus); *CCs* splenium of the corpus callosum (including posterior parietal, superior temporal, and inferior temporal/occipital subregions of the corpus callosum); *pPERI* posterior pericallosal white matter; and *Occ* occipital white matter (including forceps major)

of true effects, because nonsignificant findings were assigned an effect size of zero.

From this table, some support is present for the general trends of the anterior–posterior gradient and prominence of speed and executive functioning that we have discussed

previously. These trends do not combine in a regionally definitive manner, however, and considerable variability is evident. The largest mean effect size, for example, is in the cell representing the correlation between composite measures of executive functioning and the genu of the

corpus callosum. But several cells of the table representing the posterior regions also contain moderate effects for executive functioning and processing speed. The frontal table-cells include notable effects for other cognitive processes, including recognition memory, word reading, and postural stability.

While recognizing this variability, we would also emphasize two points: First, the vast majority of the table values are positive, reflecting a consistent relation between increasing white matter integrity and better cognitive and sensory/motor performance. Second, the selected studies were uniformly empirical in nature, but still adopted varying approaches to define the cognitive outcome measure. Whereas some studies focused on specific components of information processing (Gold et al. 2008; Madden et al. 2009), others used a broad range of tasks to construct composite measures of cognitive domains (Kennedy and Raz 2009; Zahr et al. 2009). As research in this area continues beyond these early stages, a more comprehensive account of the relation between age-related differences in white matter integrity and cognition will emerge. Using composite cognitive measures will be most valuable in investigations based on large sample sizes to establish the reliability and validity of effects, whereas using specific information processing components will be valuable in theory-driven studies that focus on the characteristics of individual neural systems (Paus et al. 2001).

Future Directions of DTI and Cognitive Aging

In this rapidly developing area of neuroimaging, progress in several directions will be valuable to the cognitive neuroscience of aging.

Clinical Application DTI is already widely used in diagnosis and prognosis of neurological disease, especially MS (Filippi et al. 2001; Ge et al. 2005; Goldberg-Zimring et al. 2005; Kealey et al. 2005), and stroke (Mukherjee 2005; Yu et al. 2009). Several studies demonstrate that DTI can provide insight into the neurobiological mechanisms of Alzheimer's disease and related dementias disease (Bozzali et al. 2001; Song et al. 2004). Empirical results from this type of application can contribute to theoretical issues in cognitive aging. For example, is there a qualitative difference between normal brain aging and dementia? Investigations that have included patients with Alzheimer's disease, as well as healthy younger and older adults, suggest that the regional changes in white matter integrity associated with dementia are not simply an exaggeration of normal aging (Damoiseaux et al. 2009; Head et al. 2004). Similarly, given the relation among cardiovascular risk factors, WMH, and cognitive decline (Rabbitt et al. 2007; van den Heuvel et al. 2006),

DTI can potentially help differentiate age-related effects from health-related effects (Raz et al. 2003, 2005, 2007).

The documentation of recovery and rehabilitation following brain injury is also an expanding area of DTI application (Johansen-Berg and Behrens 2006). Bendlin et al. (2008) tested 46 traumatic brain injury patients at two time points, approximately 2 months following their injury and 1 year later. These authors found that DTI defined areas of decreased FA and increased MD at the 1-year point, relative to controls, that were not indicated by high resolution T₁-weighted images. In response to normal adult developmental changes in gray matter and white matter, the plasticity of the central nervous system may lead to changes in cognitive strategy, which are expressed behaviorally as the age differences measured in tests of cognitive performance (Greenwood 2007). Thus, the clinical studies of brain injury may be an informative model for applying DTI to investigate the role of functional plasticity in cognitive aging.

An emerging area in neuroimaging studies of aging and functional plasticity is the effect of aerobic exercise and cardiovascular fitness. Both cross-sectional and longitudinal studies suggest that a higher level of cardiovascular fitness can help slow the trajectory of age-related decline in cortical and white matter volume (Colcombe et al. 2003; Kramer et al. 1999; Kramer and Hillman 2006). To date, studies have relied primarily on estimates of gray and white matter density, from voxel-based morphometry. Although preliminary evidence from DTI indicates that aerobic fitness is associated with increased FA, independently of age (Marks et al. 2007), additional research with DTI on this topic is necessary.

Improvements in DTI Methodology The current implementation of DTI often suffers from fluid and flow contaminations from CSF and vasculature due to partial voluming. This is particularly true in the ventral brain regions, within the periventricular space or close to the cortical surface, as a result of the hyper-intensity of fluid in the T₂ weighted baseline image (Bhagat and Beaulieu 2004). In addition, because there are many variables contributing to anisotropy within a large imaging voxel, acquisition techniques that can distinguish the various sources will be critical to understanding the mechanisms of age-related differences in diffusivity. New methodologies provide the ability to identify some of the relevant microstructural variables, such as the axon density and axon diameter distribution within a voxel (Alexander 2008; Assaf et al. 2008).

DTI involves the acquisition of diffusion-weighted images sensitized in various gradient directions. While the inhomogeneous field is the primary reason for the base image distortion, the distortion of the diffusion weighted images results from the combination of the inhomogeneous field (which is the same for the base image) and eddy currents from strong diffusion gradients. A direct method to correct for

distortion is to acquire magnetic field maps that correspond to the various diffusion weighting directions. The individual field maps can then be used to correct both the static and dynamic distortions (Chen et al. 2006). Truong et al. (2008) have recently applied this technique, combined with sensitivity encoding (SENSE) DTI acquisition, with a high-order polynomial correction to remove both distortions. Shown in Fig. 6 are colored FA maps and the original T_2^* maps. The uncorrected FA map is presented in Panel A. FA maps are presented with ΔB_0 correction (Panel B) and ΔB_0 plus eddy current correction (Panel C). Compared with the current best practice method (Panel D), a double-refocused spin-echo DTI acquisition (Reese et al. 2003), the Truong et al. method achieves further improved spatial accuracy within the frontal lobe, gray/white matter boundaries, and along the edge, as well as 19.8% higher signal-to-noise ratio.

Researchers are also actively seeking to improve the angular resolution of tensor images. Accurate reconstruction of neural connectivity patterns from DTI has been hindered by the relatively low in-plane resolution of DTI (typically 1–2.5 mm²), which limits the discrimination of crossing fibers. Techniques such as q-ball imaging (Tuch et al. 2003) and diffusion spectrum imaging (Schmahmann et al. 2007) provide estimates of multiple fiber orientations, although validation with anatomical techniques (e.g., autoradiography) has not yet been achieved (Catani 2007).

Liu and colleagues (Liu et al. 2004, 2009) have developed the self-navigated interleaved spiral (SNAILS) sequence for DTI, which can provide higher resolution than the sequences typically used in investigations of aging and cognition. SNAILS is a multi-shot, fat-saturated diffusion-weighted spin echo sequence. The technique oversamples the center of k-space, which provides an inherent motion compensation capability. Figure 7 is an example of DTI imaging acquired with SNAILS, which yielded an in-plane resolution of 390 μm^2 . This represents one of the highest spatial

resolutions that has been achieved *in vivo* on a human brain. Evident in the figure is the discrimination of multiple fiber orientations, which match the tissue morphology in corresponding myelin-stained sections.

The SNAILS and related methodologies will be particularly valuable in application to aging, because for any white matter pathway, FA is influenced by the anatomical architecture of the pathway. For example, Tuch et al. (2005), in a study of younger adults, found that increasing FA in the optic visual pathway of the right hemisphere regions was correlated with *slower* performance in a choice reaction time task, in contrast to the trends illustrated in Table 1. Tuch et al. suggest that in this case, lower FA may reflect increased crossing of fibers within the visual pathways, rather than decreased myelin integrity.

Integration with fMRI: Task-related Activation A central theme of DTI research has been the characterization of the effects of decline in white matter integrity as disconnection within the neural systems mediating cognitive functioning (Andrews-Hanna et al. 2007; Bartzokis et al. 2004; Charlton et al. 2006; Sullivan and Pfefferbaum 2006). This theoretical perspective, in turn, builds on the view that cognition is a product of distributed and interrelated neural systems (LaBerge 2000; Mesulam 1990). To develop this characterization further, in the context of aging, it will be necessary to integrate findings from DTI, behavioral measures of cognition, and functional MRI ([fMRI]; Ramnani et al. 2004; Sullivan and Pfefferbaum 2006).

An ubiquitous finding in fMRI research on cognitive aging is the increased activation for older adults, compared to younger adults, during some cognitive tasks (Dennis and Cabeza 2008; Madden et al. 2005). This age-related increased activation often occurs in frontal and parietal regions and may represent a compensatory shift to more elaborative (top-down) processing in response to age-related decline in

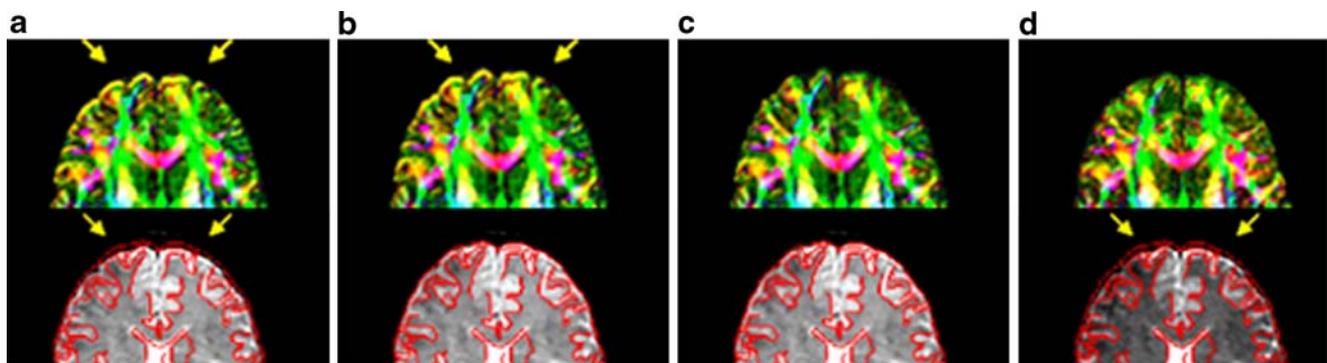


Fig. 6 FA maps (*top*) acquired with sensitivity encoding (SENSE) DTI and base image (*bottom*, with anatomical reference shown in *red*). **a** without correction; **b** with ΔB_0 correction; **c** with ΔB_0 and eddy current correction. The artificially high FA values at the edge, frontal lobe, and gray/white matter boundary are completely removed with

correction in Panel C. The technique also showed improved distortion correction and higher signal-to-noise ratio (by 19.8%), compared to the DTI acquisition using a double-refocused spin-echo sequence (**d**). Figure modified from Truong et al. (2008). © *Neuroimage* and Elsevier, 2008

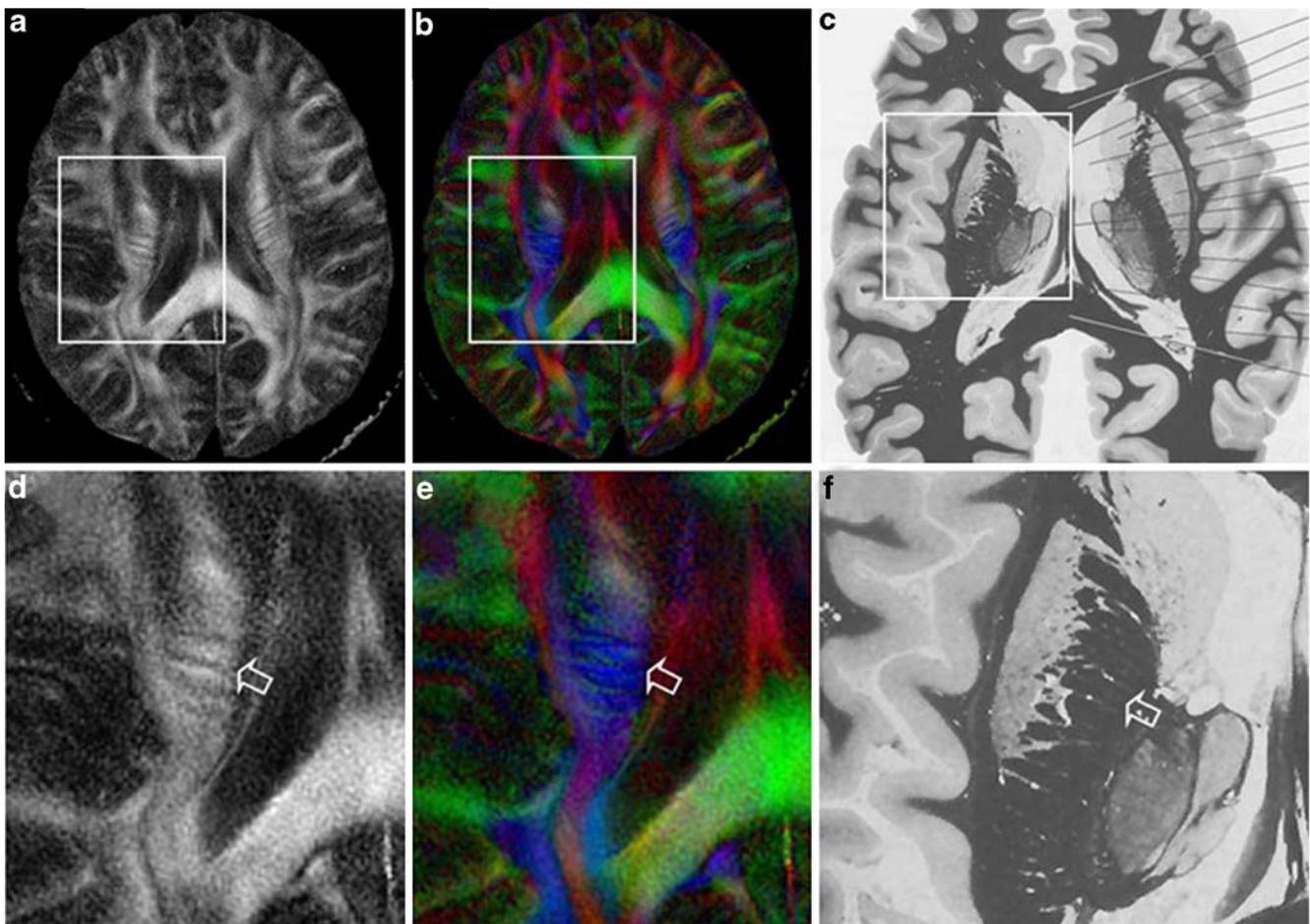


Fig. 7 High resolution self-navigated interleaved spiral (SNAILS) DTI with an in-plane resolution of $390 \times 390 \mu\text{m}^2$. **a** FA map; **b** color-coded FA map with red representing the direction of anterior–posterior, green representing the direction of left–right and blue representing the direction of superior–inferior; **c** myelin stained brain section in a similar location obtained from the Yakovlev collection (National Museum of Health and Medicine, Washington, DC); **d**, **e**,

and **f** an enlarged ROI indicated by the *rectangular boxes* in **a**, **b** and **c** showing the FA map, the color-coded FA map and the brain section respectively. Substructures of the internal capsule are shown clearly in the FA maps that match the tissue morphology (*arrow*). Figure modified from Liu et al. (2009). © *Magnetic Resonance in Medicine* and Wiley-Liss, Inc., 2009

more peripheral (bottom-up) encoding and response processing (Davis et al. 2008; Madden 2007). A related theory is that the plasticity of the aging central nervous system drives the development of behavioral strategies, with the goal of adapting to the effects of volumetric decline and related structural changes (Greenwood 2007).

An open and important question is whether age-related differences in white matter integrity have a mediating role in the age-related differences in cortical activation. Nordahl et al. (2006) found that increasing volume of prefrontal WMH, defined from structural imaging, was associated with lower levels of task-dependent fMRI activation, suggesting an interruption of the task-relevant neural systems (see also Colcombe et al. [2005]). From a compensation theory view, however, *increased* functional activation would be expected, if decreasing white matter

integrity were driving cortical recruitment. One DTI study did find that older adults with lower FA in the frontoparietal network exhibited higher fMRI activation in the superior parietal lobule, but this effect did not have a reliable role in task performance (Madden et al. 2007). An intriguing finding that is consistent with the compensation theory occurred in a study of MS patients (Rocca et al. 2007). Relative to controls, the MS patient group exhibited increased task-related fMRI activation of motor regions and increased functional connectivity between the motor regions and cerebellum. The patients' DTI data indicated that declining white matter integrity, in pathways between the motor regions and cerebellum, was associated with increased fMRI functional connectivity. Whether this type of pattern will occur in age-related comparisons of healthy individuals has yet to be determined. The Rocca et al. data are also consistent with

the superior–inferior gradient proposed by Sullivan and colleagues (Sullivan and Pfefferbaum 2006; Sullivan et al. 2008; Zahr et al. 2009).

Integration with fMRI: Intrinsic Functional Connectivity Although research interest in fMRI has concentrated on the variations in task-related activation, it is worth noting that task-related activity is only approximately 5% of the fMRI signal. The remaining 95% is spontaneous, low-frequency (< 0.1 Hz) fluctuations that reflect the intrinsic activity of the brain, which is often referred to as the default mode (Fox et al. 2005; Raichle et al. 2001). This intrinsic or default brain activity, which occurs in the absence of a specific task, also exhibits reliable patterns of regional correlation. These patterns, in turn, appear to represent the neural systems activated during cognitive tasks, such as the dorsal and ventral frontoparietal networks of attention (Fox et al. 2006). Recent DTI studies suggest that intrinsic functional connectivity depends on the structural integrity of associated white matter (Greicius et al. 2009; van den Heuvel et al. 2008).

How this intrinsic functional connectivity changes with age, and the role of white matter integrity, are largely unexplored. It is known that intrinsic functional connectivity varies with adult age. Across several investigations, intrinsic functional connectivity has been found to be lower for healthy older adults than for younger adults (Andrews-Hanna et al. 2007; Damoiseaux et al. 2008; Sambataro et al. 2008; Wu et al. 2007), with further decline evident in early-stage Alzheimer's disease (Greicius et al. 2004). Andrews-Hanna et al., conducting correlations within an older adult group, found that increasing FA, from a relatively large ROI, was associated with an increasing level of intrinsic functional connectivity.

Chen et al. (2009) developed a new method of identifying behaviorally relevant networks of intrinsic functional connectivity. These authors demonstrated that intrinsic connectivity within a prefrontal network (centered around the inferior prefrontal gyrus) was a significant mediator of the age-related variability in older adults' choice reaction time. Further, for older adults, increasing FA within the genu of the corpus callosum was correlated with increasing functional connectivity (Fig. 8). Thus, the structural and functional integrity of this prefrontal network may be a mechanism of age-related slowing of perceptual-motor speed, which in turn contributes to age-related differences in attention, memory, and other cognitive abilities (Madden 2001; Salthouse 1996; Salthouse and Madden 2007). Given that the majority of the variability in the fMRI signal reflects these spontaneous, low-frequency fluctuations, age-related differences in white matter integrity may have a more direct influence on intrinsic functional connectivity than on the task-related component of the fMRI signal.

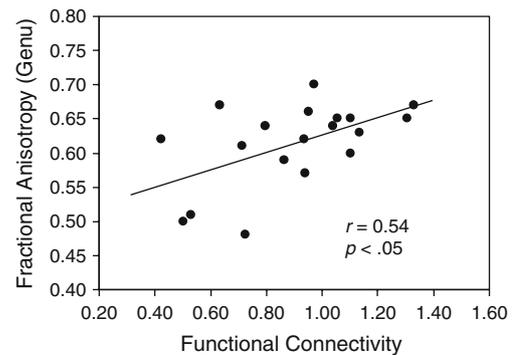


Fig. 8 Relation between white matter integrity (fractional anisotropy) in the genu of the corpus callosum and intrinsic functional connectivity (in networks connected with inferior frontal gyri), for older adults. Figure modified from Chen et al. (2009) © *Brain Structure and Function* and Springer-Verlag Berlin Heidelberg, 2009

Conclusion

The wide array of DTI research investigations has contributed significantly to the cognitive neuroscience of aging. The results confirm the role of disconnection among distributed neural systems as a fundamental mechanism of age-related variability in cognitive performance. Several trends occur across studies, notably the anterior–posterior gradient of the decline in white matter integrity with increasing age, the influence of white matter integrity on information processing speed and executive functioning, and the dependence of DTI measures on local tissue architecture. Age-related differences in white matter integrity occur throughout the brain, however, and the influence of white matter integrity has been evident in composite measures of broad functional domains, as well as in specific forms of information processing speed. To date, investigations have typically considered the relation between aging and white matter integrity, and between cognition and white matter, as separate issues. Further progress will be achieved as researchers focus on white matter integrity as a mediator of the age-related variance in cognitive performance. New information regarding the patterns of age-related differences in axial and radial diffusivity will help identify the relevant neurobiological mechanisms. These new directions in DTI research will not only help shape theories of cognitive aging but also guide translational applications to diagnosis and rehabilitation.

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References

- Abe, O., Aoki, S., Hayashi, N., Yamada, H., Kunimatsu, A., Mori, H., et al. (2002). Normal aging in the central nervous system: quantitative MR diffusion-tensor analysis. *Neurobiology of Aging*, *23*, 433–441.
- Alexander, D. C. (2008). A general framework for experiment design in diffusion MRI and its application in measuring direct tissue-microstructure features. *Magnetic Resonance in Medicine*, *60*, 439–448.
- Andrews-Hanna, J. R., Snyder, A. Z., Vincent, J. L., Lustig, C., Head, D., Raichle, M. E., et al. (2007). Disruption of large-scale brain systems in advanced aging. *Neuron*, *56*, 924–935.
- Ardekani, S., Kumar, A., Bartzokis, G., & Sinha, U. (2007). Exploratory voxel-based analysis of diffusion indices and hemispheric asymmetry in normal aging. *Magnetic Resonance Imaging*, *25*, 154–167.
- Assaf, Y., Blumenfeld-Katzir, T., Yovel, Y., & Basser, P. J. (2008). AxCaliber: a method for measuring axon diameter distribution from diffusion MRI. *Magnetic Resonance in Medicine*, *59*, 1347–1354.
- Baron, R. M., & Kenny, D. A. (1986). The moderator-mediator variable distinction in social psychological research: Conceptual, strategic, and statistical considerations. *Journal of Personality and Social Psychology*, *51*, 1173–1182.
- Bartzokis, G. (2004). Age-related myelin breakdown: a developmental model of cognitive decline and Alzheimer's disease. *Neurobiology of Aging*, *25*, 5–18.
- Bartzokis, G., Sultzer, D., Lu, P. H., Nuechterlein, K. H., Mintz, J., & Cummings, J. L. (2004). Heterogeneous age-related breakdown of white matter structural integrity: Implications for cortical “disconnection” in aging and Alzheimer's disease. *Neurobiology of Aging*, *25*, 843–851.
- Basser, P. J., & Jones, D. K. (2002). Diffusion-tensor MRI: theory, experimental design and data analysis—a technical review. *NMR in Biomedicine*, *15*, 456–467.
- Basser, P. J., Pajevic, S., Pierpaoli, C., Duda, J., & Aldroubi, A. (2000). In vivo fiber tractography using DT-MRI data. *Magnetic Resonance in Medicine*, *44*, 625–632.
- Beaulieu, C. (2002). The basis of anisotropic water diffusion in the nervous system—a technical review. *NMR in Biomedicine*, *15*, 435–455.
- Behrens, T. E., Woolrich, M. W., Jenkinson, M., Johansen-Berg, H., Nunes, R. G., Clare, S., et al. (2003). Characterization and propagation of uncertainty in diffusion-weighted MR imaging. *Magnetic Resonance in Medicine*, *50*, 1077–1088.
- Bendlin, B. B., Ries, M. L., Lazar, M., Alexander, A. L., Dempsey, R. J., Rowley, H. A., et al. (2008). Longitudinal changes in patients with traumatic brain injury assessed with diffusion-tensor and volumetric imaging. *Neuroimage*, *42*, 503–514.
- Bennett, I. J., Madden, D. J., Vaidya, C. J., Howard, J. H., Jr., & Howard, D. V. (2009). Age-related differences in multiple measures of white matter integrity: A diffusion tensor imaging study of healthy aging. *Human Brain Mapping*.
- Bhagat, Y. A., & Beaulieu, C. (2004). Diffusion anisotropy in subcortical white matter and cortical gray matter: Changes with aging and the role of CSF-suppression. *Journal of Magnetic Resonance Imaging*, *20*, 216–227.
- Bozzali, M., Franceschi, M., Falini, A., Pontesilli, S., Cercignani, M., Magnani, G., et al. (2001). Quantification of tissue damage in AD using diffusion tensor and magnetization transfer MRI. *Neurology*, *57*, 1135–1137.
- Brickman, A. M., Zimmerman, M. E., Paul, R. H., Grieve, S. M., Tate, D. F., Cohen, R. A., et al. (2006). Regional white matter and neuropsychological functioning across the adult lifespan. *Biological Psychiatry*, *60*, 444–453.
- Bucur, B., Madden, D. J., Spaniol, J., Provenzale, J. M., Cabeza, R., White, L. E., et al. (2008). Age-related slowing of memory retrieval: contributions of perceptual speed and cerebral white matter integrity. *Neurobiology of Aging*, *29*, 1070–1079.
- Burzynska, A. Z., Preuschhof, C., Bäckman, L., Nyberg, L., Li, S.-C., Lindenberger, U., et al. (2009). Age-related differences in white-matter microstructure: Region-specific patterns of diffusivity. Manuscript submitted for publication.
- Cabeza, R., Nyberg, L., & Park, D. (eds). (2005). *Cognitive neuroscience of aging: Linking cognitive and cerebral aging*. Oxford: Oxford University Press.
- Catani, M. (2006). Diffusion tensor magnetic resonance imaging tractography in cognitive disorders. *Current Opinion in Neurology*, *19*, 599–606.
- Catani, M. (2007). From hodology to function. *Brain*, *130*, 602–605.
- Catani, M., & Ffytche, D. H. (2005). The rises and falls of disconnection syndromes. *Brain*, *128*, 2224–2239.
- Catani, M., Howard, R. J., Pajevic, S., & Jones, D. K. (2002). Virtual in vivo interactive dissection of white matter fasciculi in the human brain. *Neuroimage*, *17*, 77–94.
- Charlton, R. A., Barrick, T. R., McIntyre, D. J., Shen, Y., O'Sullivan, M., Howe, F. A., et al. (2006). White matter damage on diffusion tensor imaging correlates with age-related cognitive decline. *Neurology*, *66*, 217–222.
- Charlton, R. A., Landau, S., Schiavone, F., Barrick, T. R., Clark, C. A., Markus, H. S., et al. (2008). A structural equation modeling investigation of age-related variance in executive function and DTI measured white matter damage. *Neurobiology of Aging*, *29*, 1547–1555.
- Chen, Z. G., Li, T. Q., & Hindmarsh, T. (2001). Diffusion tensor trace mapping in normal adult brain using single-shot EPI technique. A methodological study of the aging brain. *Acta Radiologica*, *42*, 447–458.
- Chen, B., Guo, H., & Song, A. W. (2006). Correction for direction-dependent distortions in diffusion tensor imaging using matched magnetic field maps. *Neuroimage*, *30*, 121–129.
- Chen, N.-K., Chou, Y.-H., & Madden, D. J. (2009). Measurement of spontaneous signal fluctuations in fMRI: Adult age differences in intrinsic functional connectivity. *Brain Structure and Function*.
- Ciccarelli, O., Catani, M., Johansen-Berg, H., Clark, C., & Thompson, A. (2008). Diffusion-based tractography in neurological disorders: concepts, applications, and future developments. *Lancet Neurology*, *7*, 715–727.
- Colcombe, S. J., Erickson, K. I., Raz, N., Webb, A. G., Cohen, N. J., McAuley, E., et al. (2003). Aerobic fitness reduces brain tissue loss in aging humans. *Journals of Gerontology. Series A, Biological Sciences and Medical Sciences*, *58*, 176–180.
- Colcombe, S. J., Kramer, A. F., Erickson, K. I., & Scalf, P. (2005). The implications of cortical recruitment and brain morphology for individual differences in inhibitory function in aging humans. *Psychology and Aging*, *20*, 363–375.
- Conturo, T. E., Lori, N. F., Cull, T. S., Akbudak, E., Snyder, A. Z., Shimony, J. S., et al. (1999). Tracking neuronal fiber pathways in the living human brain. *Proceedings of the National Academy of Sciences of the United States of America*, *96*, 10422–10427.
- Corouge, I., Fletcher, P. T., Joshi, S., Gouttard, S., & Gerig, G. (2006). Fiber tract-oriented statistics for quantitative diffusion tensor MRI analysis. *Medical Image Analysis*, *10*, 786–798.
- Correia, S., Lee, S. Y., Voorn, T., Tate, D. F., Paul, R. H., Zhang, S., et al. (2008). Quantitative tractography metrics of white matter integrity in diffusion-tensor MRI. *Neuroimage*, *42*, 568–581.
- Damoiseaux, J. S., Beckmann, C. F., Arigita, E. J., Barkhof, F., Scheltens, P., Stam, C. J., et al. (2008). Reduced resting-state brain activity in the “default network” in normal aging. *Cerebral Cortex*, *18*, 1856–1864.

- Damoiseaux, J. S., Smith, S. M., Witter, M. P., Arigita, E. J., Barkhof, F., Scheltens, P., et al. (2009). White matter tract integrity in aging and Alzheimer's disease. *Human Brain Mapping, 30*, 1051–1059.
- Dauguet, J., Peled, S., Berezovskii, V., Delzescaux, T., Warfield, S. K., Born, R., et al. (2007). Comparison of fiber tracts derived from in-vivo DTI tractography with 3D histological neural tract tracer reconstruction on a macaque brain. *Neuroimage, 37*, 530–538.
- 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.
- Davis, S. W., Dennis, N. A., Buchler, N. G., White, L. E., Madden, D. J., & Cabeza, R. (2009). Assessing the effects of age on long white matter tracts using diffusion tensor tractography. *Neuroimage, 46*, 530–541.
- Deary, I. J., Bastin, M. E., Pattie, A., Clayden, J. D., Whalley, L. J., Starr, J. M., et al. (2006). White matter integrity and cognition in childhood and old age. *Neurology, 66*, 505–512.
- DeCarli, C., Murphy, D. G., Tranh, M., Grady, C. L., Haxby, J. V., Gillette, J. A., et al. (1995). The effect of white matter hypertensity volume on brain structure, cognitive performance, and cerebral metabolism of glucose in 51 healthy adults. *Neurology, 45*, 2077–2084.
- Dennis, N. A., & Cabeza, R. (2008). Neuroimaging of healthy cognitive aging. In F. I. M. Craik & T. A. Salthouse (Eds.), *The handbook of aging and cognition* (3rd ed., pp. 1–54). New York: Psychology.
- Filippi, M., Cercignani, M., Inglese, M., Horsfield, M. A., & Comi, G. (2001). Diffusion tensor magnetic resonance imaging in multiple sclerosis. *Neurology, 56*, 304–311.
- Filley, C. M. (2005). White matter and behavioral neurology. *Annals of the New York Academy of Sciences, 1064*, 162–183.
- Fox, M. D., Snyder, A. Z., Vincent, J. L., Corbetta, M., Van Essen, D. C., & Raichle, M. E. (2005). The human brain is intrinsically organized into dynamic, anticorrelated functional networks. *Proceedings of the National Academy of Sciences of the United States of America, 102*, 9673–9678.
- Fox, M. D., Corbetta, M., Snyder, A. Z., Vincent, J. L., & Raichle, M. E. (2006). Spontaneous neuronal activity distinguishes human dorsal and ventral attention systems. *Proceedings of the National Academy of Sciences of the United States of America, 103*, 10046–10051.
- Friston, K. J., Holmes, A. P., Worsley, K. J., Poline, J. P., Frith, C. D., & Frackowiak, R. S. J. (1995). Statistical parametric maps in functional imaging: A general linear approach. *Human Brain Mapping, 2*, 189–210.
- Galvin, R. J., Heron, J. R., & Regan, D. (1977). Subclinical optic neuropathy in multiple sclerosis. *Archives of Neurology, 34*, 666–670.
- Ge, Y., Law, M., & Grossman, R. I. (2005). Applications of diffusion tensor MR imaging in multiple sclerosis. *Annals of the New York Academy of Sciences, 1064*, 202–219.
- Geschwind, N. (1965a). Disconnexion syndromes in animals and man. I. *Brain, 88*, 237–294.
- Geschwind, N. (1965b). Disconnexion syndromes in animals and man. II. *Brain, 88*, 585–644.
- Gold, B. T., Powell, D. K., Xuan, L., Jiang, Y., & Hardy, P. A. (2007). Speed of lexical decision correlates with diffusion anisotropy in left parietal and frontal white matter: evidence from diffusion tensor imaging. *Neuropsychologia, 45*, 2439–2446.
- Gold, B. T., Powell, D. K., Xuan, L., Jicha, G. A., & Smith, C. D. (2008). Age-related slowing of task switching is associated with decreased integrity of frontoparietal white matter. *Neurobiology of Aging, 29*, 100–106.
- Goldberg-Zimring, D., Mewes, A. U., Maddah, M., & Warfield, S. K. (2005). Diffusion tensor magnetic resonance imaging in multiple sclerosis. *Journal of Neuroimaging, 15*, 68S–81S.
- Grady, C. L. (2008). Cognitive neuroscience of aging. *Annals of the New York Academy of Sciences, 1124*, 127–144.
- Greenwood, P. M. (2000). The frontal aging hypothesis evaluated. *Journal of the International Neuropsychological Society, 6*, 705–726.
- Greenwood, P. M. (2007). Functional plasticity in cognitive aging: review and hypothesis. *Neuropsychology, 21*, 657–673.
- Greicius, M. D., Srivastava, G., Reiss, A. L., & Menon, V. (2004). Default-mode network activity distinguishes Alzheimer's disease from healthy aging: evidence from functional MRI. *Proceedings of the National Academy of Sciences of the United States of America, 101*, 4637–4642.
- Greicius, M. D., Supekar, K., Menon, V., & Dougherty, R. F. (2009). Resting-state functional connectivity reflects structural connectivity in the default mode network. *Cerebral Cortex, 19*, 72–78.
- Grieve, S. M., Williams, L. M., Paul, R. H., Clark, C. R., & Gordon, E. (2007). Cognitive aging, executive function, and fractional anisotropy: a diffusion tensor MR imaging study. *American Journal of Neuroradiology, 28*, 226–235.
- Gunning-Dixon, F. M., & Raz, N. (2000). The cognitive correlates of white matter abnormalities in normal aging: a quantitative review. *Neuropsychology, 14*, 224–232.
- Gunning-Dixon, F. M., & Raz, N. (2003). Neuroanatomical correlates of selected executive functions in middle-aged and older adults: a prospective MRI study. *Neuropsychologia, 41*, 1929–1941.
- Gunning-Dixon, F. M., Brickman, A. M., Cheng, J. C., & Alexopoulos, G. S. (2009). Aging of cerebral white matter: a review of MRI findings. *International Journal of Geriatric Psychiatry, 24*, 109–117.
- Guttmann, C. R., Jolesz, F. A., Kikinis, R., Killiany, R. J., Moss, M. B., Sandor, T., et al. (1998). White matter changes with normal aging. *Neurology, 50*, 972–978.
- Halligan, F. R., Reznikoff, M., Friedman, H. P., & La Rocca, N. G. (1988). Cognitive dysfunction and change in multiple sclerosis. *Journal of Clinical Psychology, 44*, 540–548.
- Head, D., Buckner, R. L., Shimony, J. S., Williams, L. E., Akbudak, E., Conturo, T. E., et al. (2004). Differential vulnerability of anterior white matter in nondemented aging with minimal acceleration in dementia of the Alzheimer type: evidence from diffusion tensor imaging. *Cerebral Cortex, 14*, 410–423.
- Holland, C. M., Smith, E. E., Csapo, I., Gurol, M. E., Brylka, D. A., Killiany, R. J., et al. (2008). Spatial distribution of white-matter hyperintensities in Alzheimer disease, cerebral amyloid angiopathy, and healthy aging. *Stroke, 39*, 1127–1133.
- Jenckens-Schinkel, A., Laboyrie, P. M., Lanser, J. B., & van der Velde, E. A. (1990). Cognition in patients with multiple sclerosis after four years. *Journal of the Neurological Sciences, 99*, 229–247.
- Jernigan, T. L., Archibald, S. L., Fennema-Notestine, C., Gamst, A. C., Stout, J. C., Bonner, J., et al. (2001). Effects of age on tissues and regions of the cerebrum and cerebellum. *Neurobiology of Aging, 22*, 581–594.
- Johansen-Berg, H., & Behrens, T. E. (2006). Just pretty pictures? What diffusion tractography can add in clinical neuroscience. *Current Opinion in Neurology, 19*, 379–385.
- Johansen-Berg, H., & Behrens, T. E. (eds). (2009). *Diffusion MRI: From quantitative measurement to In vivo neuroanatomy*. San Diego: Elsevier.
- Jones, D. K. (2008). Studying connections in the living human brain with diffusion MRI. *Cortex, 44*, 936–952.
- Kail, R. (1998). Speed of information processing in patients with multiple sclerosis. *Journal of Clinical and Experimental Neuropsychology, 20*, 98–106.
- Kealey, S. M., Kim, Y., Whiting, W. L., Madden, D. J., & Provenzale, J. M. (2005). Determination of multiple sclerosis plaque size with diffusion-tensor MR Imaging: comparison study with healthy volunteers. *Radiology, 236*, 615–620.

- Kennedy, K. M., & Raz, N. (2009). Aging white matter and cognition: differential effects of regional variations in diffusion properties on memory, executive functions, and speed. *Neuropsychologia*, *47*, 916–927.
- Keys, B. A., & White, D. A. (2000). Exploring the relationship between age, executive abilities, and psychomotor speed. *Journal of the International Neuropsychological Society*, *6*, 76–82.
- Kramer, A. F., & Hillman, C. H. (2006). Aging, physical activity, and neurocognitive function. In E. Acevedo & P. Ekeekakis (Eds.), *Psychobiology of physical activity* (pp. 45–60). Champaign: Human Kinetics.
- Kramer, A. F., Hahn, S., Cohen, N. J., Banich, M. T., McAuley, E., Harrison, C. R., et al. (1999). Ageing, fitness and neurocognitive function. *Nature*, *400*, 418–419.
- LaBerge, D. (2000). Networks of attention. In M. S. Gazzaniga (Ed.), *The new cognitive neurosciences* (2nd ed., pp. 711–723). Cambridge: MIT.
- Lawes, I. N., Barrick, T. R., Murugam, V., Spierings, N., Evans, D. R., Song, M., et al. (2008). Atlas-based segmentation of white matter tracts of the human brain using diffusion tensor tractography and comparison with classical dissection. *Neuroimage*, *39*, 62–79.
- Le Bihan, D. (2003). Looking into the functional architecture of the brain with diffusion MRI. *Nature Reviews. Neuroscience*, *4*, 469–480.
- Lindenberger, U., & Pötter, U. (1998). The complex nature of unique and shared effects in hierarchical linear regression: implications for developmental psychology. *Psychological Methods*, *3*, 218–230.
- Litvan, I., Grafman, J., Vendrell, P., & Martinez, J. M. (1988). Slowed information processing in multiple sclerosis. *Archives of Neurology*, *45*, 281–285.
- Liu, C., Bammer, R., Kim, D. H., & Moseley, M. E. (2004). Self-navigated interleaved spiral (SNAILS): application to high-resolution diffusion tensor imaging. *Magnetic Resonance in Medicine*, *52*, 1388–1396.
- Liu, C., Mang, S., & Moseley, M. E. (2009). In vivo generalized diffusion tensor imaging (GDTI) using higher-order tensors (HOT). *Magnetic Resonance in Medicine*.
- Madden, D. J. (2001). Speed and timing of behavioral processes. In J. E. Birren & K. W. Schaie (Eds.), *Handbook of the psychology of aging* (5th ed., pp. 288–312). San Diego: Academic.
- Madden, D. J. (2007). Aging and visual attention. *Current Directions in Psychological Science*, *16*, 70–74.
- Madden, D. J., Whiting, W. L., Huettel, S. A., White, L. E., MacFall, J. R., & Provenzale, J. M. (2004). Diffusion tensor imaging of adult age differences in cerebral white matter: relation to response time. *Neuroimage*, *21*, 1174–1181.
- Madden, D. J., Whiting, W. L., & Huettel, S. A. (2005). Age-related changes in neural activity during visual perception and attention. In R. Cabeza, L. Nyberg & D. Park (Eds.), *Cognitive neuroscience of aging: Linking cognitive and cerebral aging* (pp. 157–185). Oxford: Oxford University Press.
- Madden, D. J., Spaniol, J., Whiting, W. L., Bucur, B., Provenzale, J. M., Cabeza, R., et al. (2007). Adult age differences in the functional neuroanatomy of visual attention: a combined fMRI and DTI study. *Neurobiology of Aging*, *28*, 459–476.
- Madden, D. J., Spaniol, J., Costello, M. C., Bucur, B., White, L. E., Cabeza, R., et al. (2009). Cerebral white matter integrity mediates adult age differences in cognitive performance. *Journal of Cognitive Neuroscience*, *21*, 289–302.
- Malloy, P., Correia, S., Stebbins, G., & Laidlaw, D. H. (2007). Neuroimaging of white matter in aging and dementia. *The Clinical Neuropsychologist*, *21*, 73–109.
- Marks, B. L., Madden, D. J., Bucur, B., Provenzale, J. M., White, L. E., Cabeza, R., et al. (2007). Role of aerobic fitness and aging on cerebral white matter integrity. *Annals of the New York Academy of Sciences*, *1097*, 171–174.
- Mesulam, M. M. (1990). Large-scale neurocognitive networks and distributed processing for attention, language, and memory. *Annals of Neurology*, *28*, 597–613.
- Mori, S. (2007). *Introduction to diffusion tensor imaging*. Amsterdam: Elsevier.
- Mori, S., & van Zijl, P. C. (2002). Fiber tracking: principles and strategies—a technical review. *NMR in Biomedicine*, *15*, 468–480.
- Mori, S., & Zhang, J. (2006). Principles of diffusion tensor imaging and its applications to basic neuroscience research. *Neuron*, *51*, 527–539.
- Mori, S., Crain, B. J., Chacko, V. P., & van Zijl, P. C. (1999). Three-dimensional tracking of axonal projections in the brain by magnetic resonance imaging. *Annals of Neurology*, *45*, 265–269.
- Moseley, M. (2002). Diffusion tensor imaging and aging—a review. *NMR in Biomedicine*, *15*, 553–560.
- Mukherjee, P. (2005). Diffusion tensor imaging and fiber tractography in acute stroke. *Neuroimaging Clinics of North America*, *15*, 655–665.
- Nitkunan, A., Charlton, R. A., McIntyre, D. J., Barrick, T. R., Howe, F. A., & Markus, H. S. (2008). Diffusion tensor imaging and MR spectroscopy in hypertension and presumed cerebral small vessel disease. *Magnetic Resonance in Medicine*, *59*, 528–534.
- Nordahl, C. W., Ranganath, C., Yonelinas, A. P., Decarli, C., Fletcher, E., & Jagust, W. J. (2006). White matter changes compromise prefrontal cortex function in healthy elderly individuals. *Journal of Cognitive Neuroscience*, *18*, 418–429.
- Nucifora, P. G., Verma, R., Lee, S. K., & Melhem, E. R. (2007). Diffusion-tensor MR imaging and tractography: exploring brain microstructure and connectivity. *Radiology*, *245*, 367–384.
- Nusbaum, A. O., Tang, C. Y., Buchsbaum, M. S., Wei, T. C., & Atlas, S. W. (2001). Regional and global changes in cerebral diffusion with normal aging. *AJNR. American Journal of Neuroradiology*, *22*, 136–142.
- Oosterman, J. M., Sergeant, J. A., Weinstein, H. C., & Scherder, E. J. (2004). Timed executive functions and white matter in aging with and without cardiovascular risk factors. *Reviews in the Neurosciences*, *15*, 439–462.
- O’Sullivan, M., Jones, D. K., Summers, P. E., Morris, R. G., Williams, S. C., & Markus, H. S. (2001). Evidence for cortical “disconnection” as a mechanism of age-related cognitive decline. *Neurology*, *57*, 632–638.
- Parker, G. J., Haroon, H. A., & Wheeler-Kingshott, C. A. (2003). A framework for a streamline-based probabilistic index of connectivity (PICO) using a structural interpretation of MRI diffusion measurements. *Journal of Magnetic Resonance Imaging*, *18*, 242–254.
- Paus, T., Collins, D. L., Evans, A. C., Leonard, G., Pike, B., & Zijdenbos, A. (2001). Maturation of white matter in the human brain: a review of magnetic resonance studies. *Brain Research Bulletin*, *54*, 255–266.
- Peled, S. (2007). New perspectives on the sources of white matter DTI signal. *IEEE Transactions on Medical Imaging*, *26*, 1448–1455.
- Pfefferbaum, A., & Sullivan, E. V. (2003). Increased brain white matter diffusivity in normal adult aging: relationship to anisotropy and partial voluming. *Magnetic Resonance in Medicine*, *49*, 953–961.
- Pfefferbaum, A., Sullivan, E. V., Hedehus, M., Lim, K. O., Adalsteinsson, E., & Moseley, M. (2000). Age-related decline in brain white matter anisotropy measured with spatially corrected echo-planar diffusion tensor imaging. *Magnetic Resonance in Medicine*, *44*, 259–268.
- Pierpaoli, C., & Basser, P. J. (1996). Toward a quantitative assessment of diffusion anisotropy. *Magnetic Resonance in Medicine*, *36*, 893–906.
- Pierpaoli, C., Barnett, A., Pajevic, S., Chen, R., Penix, L. R., Virta, A., et al. (2001). Water diffusion changes in Wallerian degeneration

- and their dependence on white matter architecture. *Neuroimage*, *13*, 1174–1185.
- Prins, N. D., van Dijk, E. J., den Heijer, T., Vermeer, S. E., Jolles, J., Koudstaal, P. J., et al. (2005). Cerebral small-vessel disease and decline in information processing speed, executive function and memory. *Brain*, *128*, 2034–2041.
- Rabbitt, P., Scott, M., Lunn, M., Thacker, N., Lowe, C., Pendleton, N., et al. (2007). White matter lesions account for all age-related declines in speed but not in intelligence. *Neuropsychology*, *21*, 363–370.
- 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 of the United States of America*, *98*, 676–682.
- Ramnani, N., Behrens, T. E., Penny, W., & Matthews, P. M. (2004). New approaches for exploring anatomical and functional connectivity in the human brain. *Biological Psychiatry*, *56*, 613–619.
- Rao, S. M. (1995). Neuropsychology of multiple sclerosis. *Current Opinion in Neurology*, *8*, 216–220.
- Raz, N. (2000). Aging of the brain and its impact on cognitive performance: Integration of structural and functional findings. In F. I. M. Craik & T. A. Salthouse (Eds.), *Handbook of aging and cognition* (2nd ed., pp. 1–90). Mahwah: Erlbaum.
- Raz, N., Rodrigue, K. M., & Acker, J. D. (2003). Hypertension and the brain: vulnerability of the prefrontal regions and executive functions. *Behavioral Neuroscience*, *117*, 1169–1180.
- Raz, N., Lindenberger, U., Rodrigue, K. M., Kennedy, K. M., Head, D., Williamson, A., et al. (2005). Regional brain changes in aging healthy adults: general trends, individual differences and modifiers. *Cerebral Cortex*, *15*, 1676–1689.
- Raz, N., Rodrigue, K. M., Kennedy, K. M., & Acker, J. D. (2007). Vascular health and longitudinal changes in brain and cognition in middle-aged and older adults. *Neuropsychology*, *21*, 149–157.
- Reese, T. G., Heid, O., Weisskoff, R. M., & Wedeen, V. J. (2003). Reduction of eddy-current-induced distortion in diffusion MRI using a twice-refocused spin echo. *Magnetic Resonance in Medicine*, *49*, 177–182.
- Regan, D., Silver, R., & Murray, T. J. (1977). Visual acuity and contrast sensitivity in multiple sclerosis-hidden visual loss: an auxiliary diagnostic test. *Brain*, *100*, 563–579.
- Rocca, M. A., Pagani, E., Absinta, M., Valsasina, P., Falini, A., Scotti, G., et al. (2007). Altered functional and structural connectivities in patients with MS: a 3-T study. *Neurology*, *69*, 2136–2145.
- Rodriguez-Aranda, C., & Sundet, K. (2006). The frontal hypothesis of cognitive aging: factor structure and age effects on four frontal tests among healthy individuals. *Journal of Genetic Psychology*, *167*, 269–287.
- Rosenthal, R., & DiMatteo, M. R. (2001). Meta-analysis: recent developments in quantitative methods for literature reviews. *Annual Review of Psychology*, *52*, 59–82.
- Rovaris, M., Iannucci, G., Cercignani, M., Sormani, M. P., De Stefano, N., Gerevini, S., et al. (2003). Age-related changes in conventional, magnetization transfer, and diffusion-tensor MR imaging findings: study with whole-brain tissue histogram analysis. *Radiology*, *227*, 731–738.
- Salat, D. H., Kaye, J. A., & Janowsky, J. S. (1999). Prefrontal gray and white matter volumes in healthy aging and Alzheimer disease. *Archives of Neurology*, *56*, 338–344.
- Salat, D. H., Tuch, D. S., Greve, D. N., van der Kouwe, A. J., Hevelone, N. D., Zaleta, A. K., et al. (2005). Age-related alterations in white matter microstructure measured by diffusion tensor imaging. *Neurobiology of Aging*, *26*, 1215–1227.
- Salthouse, T. A. (1992). *Mechanisms of age-cognition relations in adulthood*. Hillsdale: Erlbaum.
- Salthouse, T. A. (1996). The processing-speed theory of adult age differences in cognition. *Psychological Review*, *103*, 403–428.
- Salthouse, T. A., & Madden, D. J. (2007). Information processing speed and aging. In J. Deluca & J. Kallmar (Eds.), *Information processing speed in clinical populations* (pp. 221–241). New York: Psychology.
- Salthouse, T. A., Fristoe, N., & Rhee, S. H. (1996). How localized are age-related effects on neuropsychological measures? *Neuropsychology*, *10*, 272–285.
- Salthouse, T. A., Atkinson, T. M., & Berish, D. E. (2003). Executive functioning as a potential mediator of age-related cognitive decline in normal adults. *Journal of Experimental Psychology: General*, *132*, 566–594.
- Sambataro, F., Murty, V. P., Callicott, J. H., Tan, H. Y., Das, S., Weinberger, D. R., et al. (2008). Age-related alterations in default mode network: Impact on working memory performance. *Neurobiology of Aging*.
- Schmahmann, J. D., Pandya, D. N., Wang, R., Dai, G., D'Arceuil, H. E., de Crespigny, A. J., et al. (2007). Association fibre pathways of the brain: parallel observations from diffusion spectrum imaging and autoradiography. *Brain*, *130*, 630–653.
- Schulte, T., Sullivan, E. V., Muller-Oehring, E. M., Adalsteinsson, E., & Pfefferbaum, A. (2005). Corpus callosal microstructural integrity influences interhemispheric processing: a diffusion tensor imaging study. *Cerebral Cortex*, *15*, 1384–1392.
- Shenkin, S. D., Bastin, M. E., MacGillivray, T. J., Deary, I. J., Starr, J. M., & Wardlaw, J. M. (2003). Childhood and current cognitive function in healthy 80-year-olds: a DT-MRI study. *Neuroreport*, *14*, 345–349.
- Shenkin, S. D., Bastin, M. E., Macgillivray, T. J., Deary, I. J., Starr, J. M., Rivers, C. S., et al. (2005). Cognitive correlates of cerebral white matter lesions and water diffusion tensor parameters in community-dwelling older people. *Cerebrovascular Disorders*, *20*, 310–318.
- Smith, S. M., Jenkinson, M., Woolrich, M. W., Beckmann, C. F., Behrens, T. E., Johansen-Berg, H., et al. (2004). Advances in functional and structural MR image analysis and implementation as FSL. *Neuroimage*, *23*, S208–219.
- Smith, S. M., Jenkinson, M., Johansen-Berg, H., Rueckert, D., Nichols, T. E., Mackay, C. E., et al. (2006). Tract-based spatial statistics: voxelwise analysis of multi-subject diffusion data. *Neuroimage*, *31*, 1487–1505.
- Smith, S. M., Johansen-Berg, H., Jenkinson, M., Rueckert, D., Nichols, T. E., Miller, K. L., et al. (2007). Acquisition and voxelwise analysis of multi-subject diffusion data with tract-based spatial statistics. *Nature Protocols*, *2*, 499–503.
- Song, S. K., Sun, S. W., Ramsbottom, M. J., Chang, C., Russell, J., & Cross, A. H. (2002). Dysmyelination revealed through MRI as increased radial (but unchanged axial) diffusion of water. *Neuroimage*, *17*, 1429–1436.
- Song, S. K., Kim, J. H., Lin, S. J., Brendza, R. P., & Holtzman, D. M. (2004). Diffusion tensor imaging detects age-dependent white matter changes in a transgenic mouse model with amyloid deposition. *Neurobiology of Disease*, *15*, 640–647.
- Stufflebeam, S. M., Witzel, T., Mikulski, S., Hamalainen, M. S., Temereanca, S., Barton, J. J., et al. (2008). A non-invasive method to relate the timing of neural activity to white matter microstructural integrity. *Neuroimage*, *42*, 710–716.
- Sullivan, E. V., & Pfefferbaum, A. (2006). Diffusion tensor imaging and aging. *Neuroscience and Biobehavioral Reviews*, *30*, 749–761.
- Sullivan, E. V., & Pfefferbaum, A. (2007). Neuroradiological characterization of normal adult ageing. *British Journal of Radiology*, *80*, S99–108.
- Sullivan, E. V., Adalsteinsson, E., Hedehus, M., Ju, C., Moseley, M., Lim, K. O., et al. (2001). Equivalent disruption of regional white matter microstructure in ageing healthy men and women. *Neuroreport*, *12*, 99–104.

- Sullivan, E. V., Adalsteinsson, E., & Pfefferbaum, A. (2006). Selective age-related degradation of anterior callosal fiber bundles quantified in vivo with fiber tracking. *Cerebral Cortex*, *16*, 1030–1039.
- Sullivan, E. V., Rohlfing, T., & Pfefferbaum, A. (2008). Quantitative fiber tracking of lateral and interhemispheric white matter systems in normal aging: relations to timed performance. *Neurobiology of Aging*.
- Sun, S. W., Liang, H. F., Le, T. Q., Armstrong, R. C., Cross, A. H., & Song, S. K. (2006). Differential sensitivity of in vivo and ex vivo diffusion tensor imaging to evolving optic nerve injury in mice with retinal ischemia. *Neuroimage*, *32*, 1195–1204.
- Sun, S. W., Liang, H. F., Schmidt, R. E., Cross, A. H., & Song, S. K. (2007). Selective vulnerability of cerebral white matter in a murine model of multiple sclerosis detected using diffusion tensor imaging. *Neurobiology of Disease*, *28*, 30–38.
- Thomas, C., Moya, L., Avidan, G., Humphreys, K., Jung, K. J., Peterson, M. A., et al. (2008). Reduction in white matter connectivity, revealed by diffusion tensor imaging, may account for age-related changes in face perception. *Journal of Cognitive Neuroscience*, *20*, 268–284.
- Thornton, A. E., & Raz, N. (1997). Memory impairment in multiple sclerosis: a quantitative review. *Neuropsychology*, *11*, 357–366.
- Truong, T. K., Chen, B., & Song, A. W. (2008). Integrated SENSE DTI with correction of susceptibility- and eddy current-induced geometric distortions. *Neuroimage*, *40*, 53–58.
- Tuch, D. S., Reese, T. G., Wiegell, M. R., & Wedeen, V. J. (2003). Diffusion MRI of complex neural architecture. *Neuron*, *40*, 885–895.
- Tuch, D. S., Salat, D. H., Wisco, J. J., Zaleta, A. K., Hevelone, N. D., & Rosas, H. D. (2005). Choice reaction time performance correlates with diffusion anisotropy in white matter pathways supporting visuospatial attention. *Proceedings of the National Academy of Sciences of the United States of America*, *102*, 12212–12217.
- Turken, A., Whitfield-Gabrieli, S., Bammer, R., Baldo, J. V., Dronkers, N. F., & Gabrieli, J. D. (2008). Cognitive processing speed and the structure of white matter pathways: convergent evidence from normal variation and lesion studies. *Neuroimage*, *42*, 1032–1044.
- van den Heuvel, D. M., ten Dam, V. H., de Craen, A. J., Admiraal-Behloul, F., Olofsen, H., Bollen, E. L., et al. (2006). Increase in periventricular white matter hyperintensities parallels decline in mental processing speed in a non-demented elderly population. *Journal of Neurology, Neurosurgery and Psychiatry*, *77*, 149–153.
- van den Heuvel, M., Mandl, R., Luigjes, J., & Hulshoff Pol, H. (2008). Microstructural organization of the cingulum tract and the level of default mode functional connectivity. *Journal of Neuroscience*, *28*, 10844–10851.
- Vernooij, M. W., de Groot, M., van der Lugt, A., Ikram, M. A., Krestin, G. P., Hofman, A., et al. (2008). White matter atrophy and lesion formation explain the loss of structural integrity of white matter in aging. *Neuroimage*, *43*, 470–477.
- Vernooij, M. W., Ikram, M. A., Vrooman, H. A., Wielopolski, P. A., Krestin, G. P., Hofman, A., et al. (2009). White matter microstructural integrity and cognitive function in a general elderly population. *Archives of General Psychiatry*, *66*, 545–553.
- Virta, A., Barnett, A., & Pierpaoli, C. (1999). Visualizing and characterizing white matter fiber structure and architecture in the human pyramidal tract using diffusion tensor MRI. *Magnetic Resonance Imaging*, *17*, 1121–1133.
- Wakana, S., Jiang, H., Nagae-Poetscher, L. M., van Zijl, P. C., & Mori, S. (2004). Fiber tract-based atlas of human white matter anatomy. *Radiology*, *230*, 77–87.
- Wheeler-Kingshott, C. A., & Cercignani, M. (2009). About “axial” and “radial” diffusivities. *Magnetic Resonance in Medicine*, *61*, 1255–1260.
- Wozniak, J. R., & Lim, K. O. (2006). Advances in white matter imaging: a review of in vivo magnetic resonance methodologies and their applicability to the study of development and aging. *Neuroscience and Biobehavioral Reviews*, *30*, 762–774.
- Wu, T., Zang, Y., Wang, L., Long, X., Hallett, M., Chen, Y., et al. (2007). Aging influence on functional connectivity of the motor network in the resting state. *Neuroscience Letters*, *422*, 164–168.
- Yoon, B., Shim, Y. S., Lee, K. S., Shon, Y. M., & Yang, D. W. (2008). Region-specific changes of cerebral white matter during normal aging: a diffusion-tensor analysis. *Archives of Gerontology and Geriatrics*, *47*, 129–138.
- Yu, C., Zhu, C., Zhang, Y., Chen, H., Qin, W., Wang, M., et al. (2009). A longitudinal diffusion tensor imaging study on Wallerian degeneration of corticospinal tract after motor pathway stroke. *Neuroimage*, *47*, 451–458.
- Zahr, N. M., Rohlfing, T., Pfefferbaum, A., & Sullivan, E. V. (2009). Problem solving, working memory, and motor correlates of association and commissural fiber bundles in normal aging: a quantitative fiber tracking study. *Neuroimage*, *44*, 1050–1062.
- Zhang, Y., Du, A. T., Hayasaka, S., Jahng, G. H., Hlavín, J., Zhan, W., et al. (2008). Patterns of age-related water diffusion changes in human brain by concordance and discordance analysis. *Neurobiology of Aging*.
- Ziegler, D. A., Pigué, O., Salat, D. H., Prince, K., Connally, E., & Corkin, S. (2008). Cognition in healthy aging is related to regional white matter integrity, but not cortical thickness. *Neurobiology of Aging*.