INTRODUCTION

Worldwide, the elderly population is growing, increasing the concern over age-related cognitive decline. Cognitive decline has a negative impact on quality of life and, healthy brain aging is an essential component of overall well-being. Because of the nearly global rise in life expectancy and the concomitant emphasis on quality of life, healthy brain aging has become an important focus.

Normal aging and to a greater degree degenerative brain diseases such as Alzheimer's disease (AD), cause changes in the brain's structure and function. Degenerative changes in brain structure and decline in its function are associated with declines in cognitive ability. Early detection of AD is a key priority in dementia services and research. However, depending on the disease progression, neurodegenerative manifestations, such as cerebral atrophy, are detected late in course of AD. Functional changes in the brain may be an indirect indicator of trans-synaptic activity and they usually appear prior to structural changes in AD. Resting-state functional magnetic resonance imaging (RS-fMRI) has recently been highlighted as a new technique for interrogating intrinsic functional connectivity networks. Among the majority of RS-fMRI studies, the default mode network (DMN), salience network (SN), and central executive network (CEN) gained particular focus because alterations to their functional connectivity were observed in subjects who had AD, who had mild cognitive impairment (MCI), or who were at high risk for AD. Herein, we present a review of the current research on changes in functional connectivity, as measured by RS-fMRI. We focus on the DMN, SN, and CEN to describe RS-fMRI results from three groups: normal healthy aging, MCI and AD.

Key Words Alzheimer's, Aging, Functional connectivity, fMRI, Resting state.
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imaging (RS-fMRI) has recently been highlighted as a new
technique for interrogating intrinsic functional connectivity
networks (ICN) without an experimentally determined con-
text. Previous studies have shown that ICN changes are sen-
tive to functional brain changes related to Alzheimer’s dis-
ease pathology across the clinical spectrum.

Herein, we present a review of the current research on chang-
es in functional connectivity, as measured by RS-fMRI. We
compare researcher’s outcomes from studies that examine
normal healthy aging, patients with MCI, and AD.

RESTING-STATE FMRI BASIC
PRINCIPLES AND CORE LARGE-SCALE
NETWORKS

Recently, fMRI has largely replaced nuclear medicine tech-
niques because it is non-invasive and does not require use of
radioactive contrast media, instead, fMRI uses blood-oxygen-
level-dependent (BOLD) contrast. In BOLD fMRI, the dif-
fERENCE in magnetic susceptibility between oxygenated and
deoxygenated blood serves as an intrinsic contrast medium
through which to compare conditions.

Initially, fMRI was developed to measure two conditions
(e.g., task or no task) using an activation paradigm, so-called
task-based fMRI. In recent years, there has been an increased
interest in applying the technique to at-rest conditions, termed
RS-fMRI. RS-fMRI focuses on spontaneous low-frequency
fluctuations (<0.1 Hz) in the BOLD signal. This has led to the
description of 10–20 RS networks, the most commonly
observed of which are: the default mode network (DMN), the sa-
lience network (SN), the central executive network (CEN), the
visual processing network, the sensorimotor network, the dor-
sal attention network, and auditory network, to name a few.

In many task-based fMRI studies, reversed contrasts re-
vealed consistent deactivation of specific regions, including
the medial prefrontal cortex (mPFC), the posterior cingulate
cortex (PCC), the precuneus, the anterior cingulate cortex
(ACC), the parietal cortex, and, in a minority of studies, the
hippocampus. The default mode network (DMN) is so
garded because it is the most active network in the absence of
a task. Among the majority of RS-fMRI studies, the DMN
acquired particular focus because alterations to its functional
connectivity were observed in subjects who had AD, who had
MCI, or who were at high risk for AD. The DMN is known to
be involved in the retrieval of autobiographical episodic mem-
ory as well as self-referential mental processing.

Besides the DMN, other stable ICNs, such as the SN and the
CEN, have been known to play crucial roles in cognitive pro-
cessing. The CEN includes the dorsolateral prefrontal cor-
tex (DLPFC) and the posterior parietal cortex (PPC). It plays
a key role in active maintenance and manipulation of informa-
tion in working memory, and in judgment and decision-mak-
ing in the context of goal-directed behavior. Additionally,
the SN mainly consists of the dorsal ACC and the frontoinsu-
cular cortices (FIC). It serves to identify the most relevant stim-
uli for guiding behavior in response to several internal and
external stimuli. The right FIC is a core node of the SN and
plays a critical role in switching between the CEN and the
DMN, which are known to interact competitively during
cognitive information processing. Once a salient stim-
ulus is detected, the FIC initiates the appropriate transient
control signals to engage the CEN to mediate attention, work-
ning memory and other higher-order cognitive processes, all
while disengaging the DMN. This switching mechanism
helps to focus attention on external stimuli and eventually al-
 lows the stimuli to take on added significance or saliency.

In this review, we focus on the DMN, SN, and CEN to de-
scribe RS-fMRI results from three groups: normal healthy
aging, MCI and AD.

ANALYTIC TECHNIQUES

There are three main analytic techniques used to assess
functional connectivity: the seed-based approach, indepen-
dent components analysis (ICA), and graph theory.

In the seed-based approach, the researcher selects a “seed”
of interest (region of interest, ROI) and extracts the time
course of activation at that ROI. This time course is then test-
ed for correlation with the time courses of the rest of the vox-
els of the brain to search for matches. Brain regions that are
positively correlated with each other are thought to belong to
functionally coupled networks.

In contrast, in ICA, the researcher does not need to select a
reference region. Instead, the entire set of four-dimensional
fMRI data can be decomposed into time courses and associat-
ed spatial maps that describe the temporal and spatial charac-
teristics of the data components. ICA seeks underlying com-
ponents of activation that are statistically independent of each
other. Both seed-based and ICA approaches tend to reveal the
same networks. However, seed-based analyses are sometimes
preferred because they can make group comparisons easier.

Studies using graph theory with RS-fMRI are emerging.
In graph theory, ROIs can be represented as nodes. Nodes
define anatomically homogeneous brain regions or individu-
al voxels. The fMRI data is spatially parcellated according to
a structural map of the brain, and correlations are computed
between activations in all region pairs. Typical measure-
mements taken under graph theory include average path length and global connectedness, which is the average length of the shortest connections between all pairs of nodes. Another measure of interest is the clustering coefficient, which is related to the connectedness of neighboring nodes and reflects the presence of small world structures.

RESTING-STATE FUNCTIONAL MRI ALTERATIONS IN THE NORMAL HEALTHY AGING

RS-fMRI allows for the investigation of the neural bases of age-related cognitive changes, such as attention, information processing, and working memory. To date, the DMN has been the main focus of aging-related RS-fMRI research. Many studies reported that this network’s activity during rest is necessary for memory consolidation. Additionally, amyloid deposition associated with AD is seen initially and most prominently in DMN hubs. Most of the RS-fMRI studies that focused on the DMN showed that aging is associated with decreased connectivity within this system. After assessing functional connectivity within the network, the seed-based and ICA studies produced similar results. Several studies reported reduced functional connectivity between anterior and posterior components of the DMN with advanced age. Wu et al. found greater integrity and greater preparation of DMN components in younger subjects compared to elderly subjects. The reduced RS-fMRI activity seen in the DMN in normal healthy aging subjects was correlated with executive dysfunction and decreased processing speed. Graph analysis studies further showed that decreased global connectivity and decreased nodal efficiency in several major brain hubs became more common during the aging process. Studies that focused on the modularity of functional networks reported that modularity decreased with age, indicating changes in size, composition, and the topological roles of the networks.

Age-related resting-state functional connectivity changes have been reported not only in the DMN but also in the SN and the CEN. Because there are fewer studies of the SN and the CEN, age-related resting-state functional connectivity changes in these regions are not yet well understood. Onoda et al. reported that connectivity within the SN decreased and certain internetwork connectivities (e.g., SN to auditory network, DMN to visual processing network, etc.) also decreased with aging. He et al. found age-related decreases in the positive functional connectivities between the right FIC in SN and the CEN and in the negative functional connectivities between the right FIC and the DMN, with and without atrophy correction. In a later study, He et al. also found that functional impairments of the SN may occur as early as in normal aging.

Several hypotheses have been formulated to explain the functional connectivity deficits associated with non-pathological brain aging. Age-related changes in resting state functional connectivity are thought to reflect declines in structural connectivity, especially in white matter integrity, dopaminergic deficit, and amyloid deposition in the brain.

RESTING-STATE FMRI ALTERATIONS IN PATIENTS WITH MCI AND AD

Most seed-based and ICA studies indicate that there is decreased connectivity in the DMN and other networks in subjects with MCI and AD. The majority of RS-fMRI studies found reduced DMN connectivity across widespread regions in the frontal, parietal, temporal and occipital regions of AD patients. Using graph analysis, major hubs in the brain seem to be affected by MCI and AD. Supkar et al. observed the loss of small-world organization in the functional brain network, characterized by a significantly lower clustering coefficient, which is indicative of disrupted local connectivity. Brier et al. also found that the clustering coefficient and modularity were lower in AD patients. In the same study, cognitively normal participants who harbor AD pathologies also showed reduced functional connectivity, demonstrating brain changes similar to, but smaller than, those seen in AD.

However, results regarding aberrant functions in the DMN from MCI patients were different in patterns from those of AD patients. Damoiseaux et al. reported decreased connectivity in the posterior DMN alongside increased connectivity in the ventral and anterior DMN using ICA. Also using ICA, Qi et al. found that MCI patients showed a trend toward reduced hippocampus activity compared to healthy elderly. They also found that reduced functional activity in the bilateral precuneus/posterior cingulate cortex, the right inferior parietal lobule, and the left fusiform gyrus indicated network deficiency, while increased activity in the left prefrontal cortex, the inferior parietal lobule, and the middle temporal gyrus suggested network compensation in MCI patients. Using a seed-based approach, the most recent study of MCI patients reported increased DMN connectivity between the medial prefrontal regions and the posterior cingulate and between the posterior cingulated, the parahippocampus and the anterior hippocampus. These results suggest that the DMN functions aberrantly in MCI patients, which might be due to neuronal loss in the initial phase of neurodegeneration. Increased functional connectivity between the PCC, the parahippocampus, and the anterior hippocampus in MCI patients may reflect a maladaptive mechanism.
Decreased DMN connectivity was even found in unaffected carriers of familial AD who have pathogenic presenilin-1 (PSEN1), presenilin-2 (PSEN2), or amyloid precursor protein (APP) mutations, as well as among subjects with cognitive complaints, but normal neuropsychological performance. Also, APOE ε4 carriers showed decreased connectivity of the posterior DMN and relatively increased connectivity of the SN. However, PSEN1-mutation carriers who are between 9 to 17 years old demonstrated increased functional connectivity between the posterior cingulate cortex and the medial temporal lobe regions, as well as high gray-matter volume in their temporal regions.

| Reference (year) | Intra-network | Inter-network |
|------------------|---------------|--------------|
| **Healthy aging** |               |              |
| Andrews-Hanna et al. (2007) | ↓ | NA | NA | NA |
| Damoiseaux et al. (2008) | ↓ | NA | NA | NA |
| Wu et al. (2011) | ↓ | NA | NA | NA |
| Onoda et al. (2012) | ↓ | NA | ↓ | ↓ SN-auditory, ↓ DMN-auditory |
| He et al. (2014) | ↓ | NA | ↓ | ↓ SN-CEN, ↓ SN-DMN |
| **Amyloid (+) cognitively normal subjects** | | |
| Hedden et al. (2009) | ↓ | NA | NA | NA |
| Mormino et al. (2011) | ↓ | NA | NA | NA |
| Sheline et al. (2010) | ↓ | NA | NA | NA |
| Lim et al. (2014) | ↑ | — | ↓ | ↓ DMN-CEN |
| **Mild cognitive impairment** | | |
| Damoiseaux et al. (2012) | ↓ Posterior | NA | NA | NA |
| Qi et al. (2010) | ↓ Precuneus/PCC, Rt. Inf. parietal, Lt. fusiform | NA | NA | NA |
| Gardini et al. (2015) | ↑ | NA | NA | NA |
| Sorg et al. (2007) | ↓ | NA | NA | NA |
| Agosta et al. (2012) | ↓ | NA | NA | NA |
| He et al. (2014) | NA | NA | ↑ SN-CEN, ↓ SN-DMN |
| Brier et al. (2012) | ↓ | ↓ | NA | NA |
| Wang et al. (2013) | ↓ | NA | NA | NA |
| Bai et al. (2008) | ↓ | NA | NA | NA |
| **Alzheimer's disease** | | |
| Balachandar et al. (2015) | ↓ | NA | ↑ | NA |
| Brier et al. (2012) | ↓ | ↓ | NA | NA |
| Zhang et al. (2009) | ↓ | NA | ↑ | NA |
| Agosta et al. (2012) | ↑ | ↑ | ↑ | ↑ |
| Zhou et al. (2010) | ↓ | ↑ | NA | NA |
| He et al. (2014) | NA | ↓ | ↑ SN-CEN, ↓ SN-DMN |

DMN: default mode network, SN: salience network, CEN: central executive network, ↓: decreased functional connectivity, ↑: increased functional connectivity, —: not different, NA: not applicable, PCC: posterior cingulate cortex, PFC: prefrontal cortex. RS-fMRI: Resting-state functional magnetic resonance imaging.
There are several studies that look into the association between amyloid-β deposition and functional brain connectivity. Several previous studies reported that reduced functional connectivity of the DMN can be found in cognitively normal subjects with brain amyloid burden. Using graph analysis, a decreased number of hubs and reduced connectivity within the DMN were found in both MCI and AD, as well as in healthy elderly with amyloid-β deposition on positron emission tomography (PET) images. However, Lim et al. reported that the functional connectivity of the DMN among older adults with normal cognition was greater, while the SN was not different, and the CEN was lower in among the Pittsburgh compound B-positive group, compared to the Pittsburgh compound B-negative group. The authors explained that the discrepancy in DMN result might be attributed to the “acceleration” and “brain reserve” hypotheses. The acceleration hypothesis predicts that the initiation of amyloid-β deposition by independent events instigates a milieu of higher functional connectivity that accelerates this deposition, which eventually leads to functional disconnection or metabolic deterioration in subjects with amyloid burden. The brain reserve hypothesis states that higher brain metabolism and functional connectivity represent stable traits that impart the capacity to withstand amyloid-β deposition and maintain normal cognition.

Results in the SN and CEN appear to be less consistent. Sorg et al. found reduced connectivity in the DMN and central executive network (CEN) hubs of MCI patients. Agosta et al. reported decreased connectivity in the DMN among MCI and AD patients, as well as increased mean connectivity in the CEN among AD patients. Balachandar et al. reported decreased connectivity in the DMN and increased connectivity in CEN in mild AD patients. Zhou et al. reported increased functional connectivity in the SN of AD patients. However, He et al. found that both the structural and functional organizations of the SN are impaired in MCI and AD patients. The summary of the main group differences in RS-fMRI obtained using both the seed-based and independent components analysis (ICA) methods is provided in Table 1.

CONCLUSIONS

According to a number of empirical studies, subjects who are aging healthfully, those who have MCI and those who have AD all show changes in functional connectivity in three large-scale networks—the DMN, SN, and CEN—based on RS-fMRI results. These findings help build an understanding of the neural substrates in the aging brain and during pathological degenerative processes.

Previous studies have shown that the ICN changes that are observable in RS-fMRI are sensitive to the functional brain changes that are often seen across the spectrum of AD pathology. The ICN changes seen on RS-fMRI are potentially sensitive, non-invasive biomarkers of AD. Among the ICNs, the DMN, SN and CEN have turned out to be particularly important for maintaining higher cognitive function; hence, the use of the term “core” neurocognitive networks. In particular, the DMN has been the most investigated resting-state network. And most RS-fMRI studies report reduced DMN connectivity in normally aging, MCI and AD subjects. However, reduced DMN connectivity has also been reported to be associated with a variety of other diseases, such as Parkinson’s disease and multiple sclerosis, and is by no means specific to AD. Besides, DMN, SN and CEN abnormalities are widespread in psychiatric, as well as in neurological disorders. Different studies have revealed that there are both consistencies and discrepancies in ICN changes among normally aging, MCI and AD subjects, possibly due to methodological differences and/or clinical heterogeneity among subjects. These limitations have recently been overcome and it is expected that clinical applications of RS-fMRI in normal aging, MCI and AD will continue to increase.

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