Default Mode Network Disruption Secondary to a Lesion in the Anterior Thalamus

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Objective: To describe the neuroanatomical correlations of an isolated lesion in the anterior thalamus using functional imaging in a 40-year-old man with multiple sclerosis.

Design: Case report with 10 cognitively normal controls.

Setting: Mayo Clinic, Rochester, Minnesota.

Patient: A 40-year-old man with a 2-week course of acute-onset amnesia, abulia, poor concentration, hypersomnolence, and reclusiveness.

Intervention: Functional magnetic resonance imaging.

Results: Magnetic resonance imaging demonstrated a large gadolinium-enhancing plaque in the left anterior thalamus and other demyelinating plaques in the subcortical and periventricular white matter, consistent with the diagnosis of multiple sclerosis. His symptoms persisted at the 7-month follow-up. The patient's resting state functional magnetic resonance image demonstrated an asymmetric disruption of the posterior cingulate portion of the default mode network ipsilateral to the left thalamic lesion.

Conclusions: A large multiple sclerosis plaque in the deep gray matter altered the resting state functional connectivity in a patient presenting with pure cognitive dysfunction. Such altered connectivity may underlie cognitive symptoms in neurologic disease. In addition, this case provides lesional evidence of default mode network circuitry involving the pathways of the circuit of Papez.


IN 1937, PAPEZ1 DESCRIBED A CLASSICAL CIRCUIT IMPORTANT IN HUMAN MEMORY AND EMOTION. ANATOMICALLY, THIS CIRCUIT INCLUDES PROJECTIONS FROM THE SUBICULAR COMPLEX TO THE MAMILLARY NUCLEI THAT PROJECT TO THE ANTERIOR NUCLEUS OF THE THALAMUS. THE ANTERIOR NUCLEUS OF THE THALAMUS, IN TURN, PROJECTS TO THE POSTERIOR CINGULATE CORTEX BACK TO THE HIPPOCAMPAL FORMATION. IF THIS LOOP IS DISRUPTED, SEVERE AMNESIA MAY RESULT.

FUNCTIONAL MAGNETIC RESONANCE IMAGING (fMRI) ANALYSIS TECHNIQUES USING LOW-FREQUENCY FLUCTUATIONS (0.08-0.01 HZ) IN THE BLOOD OXYGEN LEVEL–DEPENDENT (BOLD) T2* SIGNAL HAVE CONSISTENTLY IDENTIFIED GLOBAL BRAIN NETWORKS THAT ARE ANATOMICALLY AND FUNCTIONALLY RELATED.2-4 MOST OF THESE INVESTIGATIONS HAVE EXAMINED BOLD SIGNAL FLUCTUATIONS IN THE RESTING BRAIN, AND THE SUBSEQUENTLY IDENTIFIED NETWORKS HAVE BEEN TERMED RESTING STATE NETWORKS (RSN). THE RELATIONSHIP OF RSN TO COGNITIVE FUNCTION IS UNCERTAIN; IF THEY ARE RELATED TO NORMAL COGNITIVE FUNCTION, THEN IT WOULD BE REASONABLE TO HYPOTHESIZE THAT AN ACUTE CHANGE IN COGNITIVE FUNCTION WOULD LEAD TO A MEASURABLE CHANGE IN A PARTICULAR RSN OR ITS RELATIONSHIP TO OTHER RSNs.

WE ATTEMPT TO VALIDATE THIS LINE OF REASONING USING AN ILLUSTRATIVE CASE STUDY OF ACUTE COGNITIVE DYSFUNCTION ASSOCIATED WITH A LESION IN THE ANATOMICAL CIRCUIT OF PAPEZ, ANALYZED USING A RESTING-STATE fMRI METHOD KNOWN AS INDEPENDENT COMPONENT ANALYSIS (ICA). INDEPENDENT COMPONENT ANALYSIS IS A MODEL-FREE, DATA-DRIVEN fMRI ANALYSIS TECHNIQUE THAT CAN BE USED TO DEFINE THE SPATIAL AND TEMPORAL RELATIONSHIP OF LOW-FREQUENCY BOLD FLUCTUATIONS ACROSS THE RESTING BRAIN TO IDENTIFY RSN.5 IN ADDITION, WE CONDUCTED A CONNECTIVITY ANALYSIS USING CORRELATIONS OF THE BOLD TIME COURSE DERIVED FROM A SEED APPROXIMATING THE PATIENT'S LESION IN THE ANTERIOR THALAMUS.

THE DEFAULT MODE NETWORK (DMN) IS AN RSN COMPOSED OF MULTIPLE BRAIN REGIONS THAT ARE MORE ACTIVE DURING PASSIVE REST THAN DURING ATTENTION-DEMANDING TASKS.4 THE DMN IS COMPOSED OF THE POS-
terior cingulate cortex (PCC), medial prefrontal cortex, inferior parietal lobe, lateral temporal cortex, and hippocampal formation.\(^7\) The functional circuits that underlie the coordination of these diverse brain structures are currently unknown. However, these structures are within limbic, paralimbic, or heteromodal association cortices, and we hypothesize that they are interconnected via anterior and posterior limbic circuitry. The posterior limbic circuitry has commonly been referred to as the circuit of Papez,\(^1\) and its disruption has been implicated in the cognitive dysfunction present in conditions such as Wernicke-Korsakoff syndrome,\(^8\) focal hemorrhage,\(^9\) and mesial temporal sclerosis.\(^10\) In this article we use resting-state fMRI to compare the DMN activity of 10 healthy control subjects with that of a 40-year-old man with an acute lesion in the circuit of Papez leading to cognitive dysfunction.

**REPORT OF A CASE**

A 40-year-old, right-handed man experienced acute onset of personality change, hypersonomnolenece, and cognitive decline marked by poor concentration. He began sleeping more than 12 hours per day. He forgot how to operate machinery at work and became carefree and mellow.

Ten years earlier, he had an isolated 2-week episode of foot numbness. He smoked cigarettes daily but otherwise had no relevant medical history and took no medications.

At the time of referral, cognitive symptoms had been present for 2 weeks. He scored 22 of a 38 possible points on the short test of mental status,\(^11\) losing points for orientation (score, 3 of a possible 8), calculation (2 of 4), abstraction (0 of 3), attention (5 of 7), learning (3 of 4), information (3 of 4), and recall (2 of 4). He had global cognitive dysfunction, and language was impaired. He was especially poor at naming (correctly identifying 2 of 4 objects), describing a picture, and writing. He had difficulty operating machinery at work and became carefree and mellow.

Resting-state gradient, echo-planar BOLD (fMRI) sequences were obtained for the patient 12 days after initial presentation while he was still symptomatic using a General Electric 3T Signa HDx (Waukesha, Wisconsin) scanner at 14.0 level software (8-channel head coil; 30 axial slices; thickness, 5 mm without gap; time to repetition, 2000 milliseconds; echo time, 30 milliseconds; flip angle, 90\(^\circ\); in-plane resolution, 64 \times 64; field of view, 220 \times 220 mm; 200 volumes), and a spoiled gradient recalled T1-weighted structural image was obtained for coregistration of functional data. Ten cognitively normal volunteers (mean [SD] age, 44.2 [9.16] years; 3 male) were evaluated with the same imaging protocol for comparison. All participants were instructed to rest comfortably in the scanner with their eyes open. The images were preprocessed using statistical parametric mapping (SPM) software (Wellcome Department of Cognitive Neurology, University College London, London, England), with the SPM5 version implemented in MATLAB (Mathworks Inc, Sherborn, Massachusetts). Preprocessing steps included discarding the first 10 volumes to allow for scanner and cognitive equilibrium, rigid body motion correction, slice timing correction, normalization to Montreal Neurologic Institute standard space, coregistration to spoiled gradient recalled T1-weighted structural image, and smoothing with an 8-mm full width at half maximum Gaussian filter. These images were then analyzed using Group ICA of fMRI Toolbox software with individual subject maps produced with the dual-regression method.\(^12\) The independent components were visually inspected to identify the DMN based on its typical appearance. The DMN for the subject was then overlaid on the coregistered structural MRI, and the group DMN was overlaid on a representative control subject's coregistered structural MRI. To assess variability in the individual DMN components produced, the average z score from each subject's component were extracted from right and left seed regions using the MarsBar toolbox (http://marsbar.sourceforge.net/). The seed regions of interest were defined using the precuneus and PCC regions from the automated anatomical labeling atlas.\(^13\) To quantitatively compare the patient's DMN with the control group's, the 2-sample t test function in SPM5 was used in a method similar to the commonly used clinical comparison of individual subjects' positron emission tomographic scans to control groups.\(^14\) Result are reported as significant if they exceed a voxel-level false discovery rate corrected P < .01 with a corrected cluster level P < .001. A similar statistical comparison was performed for multiple other easily identifiable RSN in this data set (i.e., right central executive, left central executive, visual, auditory, and sensor-motor networks).

To further investigate the relationship of the thalamus to RSNs, a seed-based connectivity analysis was performed using the resting-state fMRI data analysis toolkit (http://www.restfmri.net), SPM5, and in-house developed software implemented in MATLAB. Prior to performing seed analysis, the resting-state images were put through several additional standard preprocessing steps that included linear detrending to correct for signal drift, 0.01- to 0.08-Hz bandpass filtering to reduce non-neuronal contributions to BOLD fluctuations, and regression correction for spurious variables such as rigid body transformation motion effects, global mean signal, and white matter and cerebral spinal fluid signal.\(^15\) Then, a 3-mm radius seed region of interest (Montreal Neurological Institute coordinates, \(-8, -9, -1\)) approximating the position of the subject's thalamic lesion was used to extract the BOLD time course for each
participant. The low-frequency time course for each participant was then correlated with every other voxel in the brain using the Pearson correlation coefficient. Regions with a fluctuating time course in phase with the region of interest time course have positive correlation values, and regions with out-of-phase fluctuations have negative correlation values. To evaluate if regions displayed a 90° out-of-phase lagging relationship (which would have a correlation value near 0), a lagged correlation analysis was performed by shifting the region of interest reference time course for each subject by a lag correspond-
ing to 90° of the reference time-course period and repeating the correlation analysis. This lagged correlation approach may have the potential to elucidate a temporal lag in a functional relationship along a circuit.16 After performing a Fischer r-to-z transformation for all images, the subject's lagged voxel-wise connectivity map was compared with those of controls (given that the lagged correlational map was consistent with 2 key DMN regions). A similar SPM statistical procedure used for the ICA analysis was used for the lagged correlation comparison. Results were reported as significant if they exceed a voxel-level false discover rate corrected $P < .01$ with a corrected cluster level $P < .001$.

**RESULTS**

The patient's ICA-identified DMN demonstrated an asymmetric disruption of the PCC portion of the default mode network ipsilateral to the left thalamic lesion (Figure 2, A). The group ICA of the resting state MRI produced the typical symmetric spatial extent of the DMN (Figure 2, B). None of the control subjects' individual DMNs displayed right to left asymmetry. The right and left PCC/precuneus contribution to every participant's DMN identified using ICA is displayed in a box plot (Figure 3). Statistical analysis of the patient's ICA-identified DMN compared with that of the control group demonstrated a reduction in the patient's DMN connectivity in the PCC, precuneus, and left inferior parietal lobe (Figure 4, A). No statistically significant difference was found in any other RSN examined.

The anterior thalamus seed-based connectivity analysis demonstrated significant group correlations (Figure 5) and patient to group comparison (Figure 4 and Figure 5). The group in-phase relationship mainly consisted of bilateral thalamic regions and regions of the saliency network (ie, bilateral frontoinsular cortices and anterior cingulate), with some additional in-phase relationship with the PCC. After time-lagged correlation, the relationship no longer revealed any thalamic contribution (as expected) but did reveal a relationship with 2 key DMN nodes (the precuneus and medial prefrontal cortex). The out-of-phase relationship consisted primarily of sensorimotor regions and the orbitofrontal cortex.

The statistical analysis of the patient's time-lagged correlational relationship compared with that of the group revealed a similar decrease in DMN regions, as was observed for the ICA analysis, which included the PCC and precuneus (Figure 4 and Figure 5).

**COMMENT**

The index case and resting state fMRI analysis illustrate 2 key points. First, deep gray matter lesions may contribute to cognitive dysfunction in neurologic illness. Second, anterior thalamic lesions along the circuit of Pa-
pez disrupt the DMN. The DMN is the brain state that dominates in a resting non–task-directed mode of brain function. To perform task-directed activities, the activity of the DMN subsides as task-relevant regions of the brain predominate. Another RSN known as the saliency network has been theorized to facilitate the switching of brain states from the DMN to task-relevant networks. The patient that we describe presented with a lesion in the thalamus that disrupted the DMN and potentially impaired his ability to switch from a default mode of brain function to a task-related state of brain function. This manifested clinically as acute cognitive dysfunction in multiple domains and persistent short-term memory impairment. Therefore, disruption of anatomical connections also manifest altered RSN functional connectivity.

This case study supports the conclusion that RSN are related to cognitive function and have the potential to advance the understanding of cognitive dysfunction in clinical settings. In addition, this case provides lesional evidence of DMN circuitry involving the pathways of the circuit of Papez. A previous positron emission tomography study has demonstrated a similar impairment in PCC function with circuit of Papez lesions secondary to alcoholic Korsakoff syndrome. However, resting-state fMRI DMN analysis provides a novel noninvasive means of investigating the relationship between anatomic lesions and cognitive symptoms.

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REFERENCES


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