Impaired Brain Pathways May Cause Attention Problems after Stroke

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OAK BROOK, Ill. — Damage to some of the pathways that carry information throughout the brain may be responsible for attention deficits in patients who have had a subcortical stroke in the brain’s right hemisphere, according to a study published online in the journal *Radiology*. Researchers hope the findings may provide a measure for selecting suitable patients for early interventions aimed at reducing cognitive decline following stroke.

A stroke may affect cortical regions of the cerebral cortex, which includes the gray matter that lines the surface of the brain, or it may affect brain regions below the cortex, including white matter tracts connecting different regions of the brain. A stroke affecting brain structures below the cortex is known as a subcortical stroke.

More than one-third of patients experience cognitive decline after a stroke, including attention deficit, which can affect and impair the patient’s ability to carry out routine activities of daily living.

“Impairment of attention has been observed in patients with both cortical and subcortical stroke,” said senior study author Chunshui Yu, M.D., from the Department of Radiology at Tianjin Medical University General Hospital in Tianjin, China. “In cortical stroke, the direct involvement of cortical regions associated with attention may account for the deficit. However, the parts of the nervous and brain systems underlying attention deficit in subcortical stroke remain largely unknown.”

To investigate the mechanisms underlying attention deficit in chronic subcortical stroke, Dr. Yu and colleagues combined voxel-based lesion-symptom mapping (VLSM) and diffusion tensor tractography (DTT) in 49 patients (32 men and 17 women between the ages of 40 and 71) after subcortical stroke and 52 control patients (30 men and 22 women, age 40-68). VLSM is a method of analyzing relationships between tissue damage and behavioral deficits, and DTT is an MRI technique that allows for 3-D visualization of specific white matter tracts in the brain.

A modified version of the attention network test was used to assess visual attention function. VLSM was used to identify lesion locations related to attention deficit in the stroke patients. Then DTT was used to determine the responsible impaired brain connections at the chronic stage (> 6 months post-stroke).

The results showed that compared to the controls, patients with chronic stroke exhibited prolonged reaction time during the attention task. VLSM revealed that having an acute stroke lesion in the right caudate nucleus and nearby white matter was correlated to the prolonged reaction time. DTT showed that the responsible lesion was located in the right thalamo- and caudate-prefrontal pathways in controls.

The right brain damage subgroup had significantly decreased fractional anisotropy (FA) in these pathways, which were correlated with the prolonged reaction time. FA provides a way to measure diffusion occurring within a region of the brain. FA is typically higher in brain regions of high organization. Reductions in FA have been previously associated with advancing age and in cases of cognitive impairment.

“The impairment of the right thalamic- and caudate-prefrontal pathways was consistently associated with attention deficit in patients with right subcortical stroke,” Dr. Yu said. “Based on this association, one can estimate which patients with stroke would be more likely to develop into long-term persisting attention deficit by evaluating the lesion-induced damage to these pathways.”

At A Glance

- A stroke affecting brain structures below the cortex is known as a subcortical stroke.
- Impairment of specific brain pathways was consistently associated with attention deficit in patients with right subcortical stroke.
- More than one-third of patients experience cognitive decline after a stroke, including attention deficit, which can affect and impair the patient’s ability to carry out routine daily activities.

“Connection Disruption Underlying Attention Deficit in Subcortical Stroke.” Collaborating with Dr. Yu were Jingchun Liu, M.S., Caihong Wang, M.S., Qingqing Diao, M.S., Wen Qin, Ph.D., and Jingliang Cheng, Ph.D.

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Figure 1. Flow diagram shows the study participants. DTI = diffusion-tensor imaging, VLSM = voxel-based lesion-symptom mapping.

High-res (TIF) version
(Right-click and Save As)
Figure 2. Flow diagram of the analysis. AAL = automated anatomic labeling, ANT = attention network test, FA = fractional anisotropy, RT = reaction time, VLSM = voxel-based lesion-symptom mapping.

Figure 3. Stroke lesions. A, Lesion incidence map in patients with acute stroke. B, Lesion incidence map shows regions in which at least 10 patients had a lesion. Color bar denotes the probability of lesion distribution. C, Brain region that is correlated with attention deficit in the voxel-based lesion-symptom mapping (VLSM) analysis. Color bar denotes the t values.
Figure 4. Anatomic connections of the responsible lesion region (red) and the thalamic-prefrontal pathway (blue). Green denotes the seed region (responsible lesion region), gray indicates the origin (thalamus) and target (prefrontal) regions of the thalamic-prefrontal pathway.
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Figure 5. Average normalized connection probabilities and fingerprints. A, Average normalized connection strengths between the entire responsible lesion region and the 44 target brain regions in the same hemisphere. B, Connection fingerprints of the entire responsible lesion region. C, Average normalized connection strengths between the involved right caudate nuclei and the 44 target brain regions in the same hemisphere. D, Fingerprints of the involved right caudate nuclei.

Figure 6. Fractional anisotropy (FA) analyses of the responsible fiber pathways. FA difference between patients with right-sided brain damage (RBD) and control subjects in the, A, right thalamic-prefrontal and, C, caudate-prefrontal pathways. Correlations between FA values of the, B, right thalamic-prefrontal and, D, caudate-prefrontal pathways and reaction time of correct responses in patients with RBD.