Reduced cerebral blood flow in white matter in ischaemic leukoaraiosis demonstrated using quantitative exogenous contrast based perfusion MRI

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Abstract

Objective—White matter hypoperfusion may play a part in the pathogenesis of ischaemic leukoaraiosis, but demonstration of this requires a high resolution quantitative method of cerebral blood flow (CBF) measurement. Initial exogenous contrast based MRI methods only allowed measurement of relative cerebral blood volume (CBV) values, but more recently a mathematical approach has been developed which enables absolute regional CBF and CBV to be determined. This technique was applied to patients with ischaemic leukoaraiosis to determine whether reduced white matter CBF in this patient group could be demonstrated.

Methods—Eight patients with ischaemic leukoaraiosis (radiological leukoaraiosis and clinical lacunar stroke), and nine age matched controls were studied. A spin echo echoplanar image sequence was used on a 1.5 Tesla MR system. An arterial input function was obtained from voxels placed over the middle cerebral arteries. Cerebral blood flow, CBV, and mean transit (MTT) maps were derived. Regions of interest were placed at standard positions in the white and grey matter and mean values of CBF, CBV, and MTT were compared between the two groups.

Results—Mean (SD) white matter CBF was significantly reduced in patients by 38% (13.40 (4.87) v 21.74 (3.53) ml/min/100 g, p=0.002). Significant reductions in CBF were seen in all white matter regions. By contrast there was no reduction in CBF in any grey matter region. There was no significant difference in white matter CBF between cases and controls; mean values were lower in all white matter regions for patients but this did not reach significance for any region. By contrast mean grey matter CBV was significantly higher in patients than in controls. Mean MTT values were higher in all regions of grey and white matter in the patient group, but this only achieved significance for the superior white matter.

Conclusion—A quantitative MR perfusion method showed reduced white matter CBF in patients with ischaemic leukoaraiosis, but normal grey matter CBF. This is consistent with hypoperfusion playing a part in the pathogenesis of ischaemic leukoaraiosis. The absolute values of white matter and grey matter CBF obtained in the patient groups were very similar to those in previous PET studies, providing further evidence for the validity of the regional CBF measurements obtained using this quantitative MR perfusion technique. The high spatial resolution and lack of radioactive administration makes such techniques ideal for longitudinal studies in this condition.

Keywords: magnetic resonance imaging; cerebrovascular circulation; stroke; cerebral blood flow.

Stroke is the third most common cause of mortality and the major cause of disability in western countries. Up to 25% of strokes are caused by lacunar infarction which results from ischaemia in the territory of the small perforating intracerebral arteries. A subgroup of these patients may have multiple lacunar infarcts which, in addition to recurrent lacunar stroke, may present with gait disturbance, a subcortical dementia, and a parkinsonian syndrome. In addition to focal lacunar infarction, subcortical vascular disease may result in diffuse white matter changes characterised by neuronal loss, demyelination, and gliosis. Its radiological appearance is termed leukoaraiosis, in view of the periventricular lucency seen on CT and a corresponding high signal on T2 MRI. Post-mortem studies show that severe leukoaraiosis is almost always accompanied by multiple lacunar infarcts, and the small perforating arteries seem abnormal with hyalination and thickening of the vessel wall. Some features suggest that leukoaraiosis results from chronic ischaemic damage. A striking feature is that the white matter changes spare the subcortical U fibres. This is consistent with the pathology being maximal in the regions of terminal supply of the penetrating arterioles—namely, the periventricular regions and the centrum semiovale. These areas, where perfusion pressure will be lowest, are likely to be particularly vulnerable to periods of relative hypoperfusion.

For these reasons investigators have attempted to show reduced perfusion or cerebral blood flow (CBF) in the white matter in patients with ischaemic leukoaraiosis. Such studies require a quantitative estimate of cerebral blood flow, as the disease process is diffuse and may affect multiple brain regions.
Semiquantitative techniques such as single photon emission computed tomography (SPECT) have been used, but these can only give ratios of perfusion between different brain regions, and cannot demonstrate a reduction in absolute regional CBF. Quantitative xenon-CT and PET have shown reduced cerebral blood flow in patients with ischaemic leukoaraiosis, but such techniques require exposure to radiation, making them unsuitable for serial imaging studies.

Recent advances in ultrafast MRI now offer the potential of combining high resolution structural imaging of the white matter with perfusion imaging. Perfusion can be determined with MRI by the use of bolus tracking dynamic imaging methods with exogenous paramagnetic contrast agents. Initial perfusion studies only allowed measurement of relative CBV values, but more recently, a mathematically advanced approach has been developed to also yield relative CBF. After dose normalisation of the gadolinium contrast agent, and introduction of a proportionality constant derived from studies in a pig model and in humans, absolute regional CBF can be calculated from MRI with a signal to noise and spatial resolution equal or better than that obtained with PET. In comparisons between MRI and PET, performed both in pigs and in healthy humans, a close correlation was found between CBF measurements calculated using the two methods.

In this study we examined the hypothesis that reduced white matter perfusion could be shown in patients with ischaemic leukoaraiosis using quantitative exogenous contrast based perfusion MRI. We determined whether the absolute values of CBF obtained were similar to those from previous studies using PET. We also compared regional cerebral blood volume (CBV) and mean transit time (MTT) between patients and age matched controls.

**Subjects and methods**

**Subjects**

We studied eight patients (seven men) presenting to a specialised cerebrovascular neurology clinic with ischaemic leukoaraiosis. This was defined as extensive confluent areas of hyperintensity in white matter on T2 weighted images, in combination with a history of lacunar stroke. All patients were studied at least 3 months after their last stroke to reduce an influence on cerebral perfusion patterns from diaschisis after an acute event. Mean time since last presentation was 2.9 years (range 1–7 years). Treated hypertension was present in six and five were ex-smokers or current smokers. None had diabetes. Five had evidence of mild cognitive impairment on routine neuropsychological testing. Three had had only single clinical lacunar strokes and the remainder had experienced recurrent events. Carotid stenosis (>30%) was excluded in all by duplex ultrasound. In addition, we studied nine age matched normal controls (eight men) of whom two had treated hypertension, but none had diabetes. These were recruited from a community volunteer database from the same geographical region as the patients. Patients with a history of stroke, transient ischaemic attack, or cognitive impairment were excluded but patients with vascular risk factors were included. Six were current or ex-smokers. Mean (SD) age was 63.3 (12.3) in patients and 68.7 (7.3) in controls (p=0.300). All patients gave informed signed consent and the study was approved by the local hospital ethics committee.

**MR imaging**

A 1.5T GE Signa (GE Medical Systems, Milwaukee, WI, USA) system fitted with Advanced NMR (Wilmington, MA, USA) hardware and software was used. Before subjects were placed in the magnet in the supine position a 16 gauge cannula was inserted into an antecubital vein. In six patients and all controls end tidal carbon dioxide monitoring was performed throughout the procedure; a nose clip was fitted to ensure that the inhaled gas was derived solely from a mouthpiece which was in place throughout the scan, and was fitted with a one way valve to separate inspired and expired gases.

An 11 slice SPGR sagittal localiser image was acquired (TR=150 ms, TE=42 ms, a=90°, Δx=Δy=0.94 mm, slice thickness=5.0 mm, slice gap=2.5 mm). The localiser image was displayed and used to prescribe image planes for the perfusion study. Five axial slices were prescribed though the region of interest, with the centre of the second uppermost slice aligned with the superior aspect of the corpus callosum. Two additional near axial oblique slices were prescribed to ensure that one intersected the middle cerebral arteries (MCAs). To check that one of the lower slices intersected at least one of the MCAs, a single seven plane spin echo (SE) echoplanar image (EPI) was acquired (TR=1.52 ms, TE=90 ms, a=90°, Δx=Δy=1.6 mm, field of view 400×200 mm, slice thickness=5.0 mm, slice gap=2.0 mm). This procedure was repeated if necessary, with the two oblique slices repositioned, until the MCAs were clearly seen within one of the slices.

Next, the subject was asked to close his or her eyes, and a series of 70 consecutive multiplanar SE-EPIs were acquired (dummy scans=4, TR=1.52 ms, TE=90 ms, a=90°, Δx=Δy=1.6 mm, slice thickness=5.0 mm, slice gap=2.0 mm) with the subject breathing air. During image acquisition, end tidal CO2 measurements were recorded. Contrast agent (Omniscan, Nycomed Imaging) at a dose of 0.1 mmol kg⁻¹ was delivered at a rate of 5 ml/s using a MR compatible power injector (Doltron), commencing on the 20th image in the series. Contrast agent administration was followed with a 10–20 ml saline flush, depending on the weight of the subject.

The image containing the MCAs was displayed as a ΔR, map, with ΔR, calculated using the first image in the series as a reference. Voxels in the vicinity of both MCAs, with large signal change and rapid bolus passage, were selected to produce a mean arterial input function (AIF). After spatial smoothing with a
uniform 3×3 convolution filter the tissue ΔR₂
time curve was deconvolved with the AIF.
Maps of CBF were derived from the height of
the deconvolved impulse response, and abso-
lute CBF was calculated by normalisation to
the injected dose multiplied by an empirical
constant factor. Deconvolution was performed
between the AIF arrival time, and the point at
which the AIF ΔR₂ time curve returned to
the noise level. The CBV was calculated pixel
by pixel as the time integral of the tissue ΔR₂
time curve. Integration was performed numerically
between the arrival time of the AIF, and the
time the ΔR₂ time curve returned close to
baseline levels, but before tracer recirculation
became apparent. Maps of MTT derived using
the central volume theorem from MTT=CBV/
CBF. No correction for the effect of delay in
arrival time in successively acquired slices or
due to acquisition order was made as previous
simulations have shown such errors to be small
at the flow values measured in this study.⁹

A T2 weighted image was created from the
mean of the precontrast SE-EPI images. Re-
gions of interest (ROIs) were defined separately
for the left and right hemispheres in the five
superior slices for the following brain regions:
frontal cortex, parietal cortex, occipital cortex,
basal ganglia, anterior periventricular white
matter; posterior periventricular white matter;
and the white matter superior to the ventricles.
The ROIs were hand drawn on the EPI T2
weighted images (mean of the prebolus images)
by an experienced observer. The ROIs were
transferred to the CBF images for analysis.
Standard brain regions, as outlined above, which
had previously been drawn on a normal MR
template image were used. After the ROIs were
transferred to the CBF images, they were hand
tored to eliminate any voxels within the vicinity
of large vessels, which show up as regions of
extremely high intensity on the CBF maps.
Weighted mean CBF, CBV, and MTT values
were calculated from the ROIs, for each brain
structure separately as

\[ P = k \cdot \frac{\sum I_1 \cdot A_1}{A_1} \]

where \( P \) is each of CBF, CBV, and MTT. \( A \)
is the area of region, \( I_1 \) its mean intensity, and \( k \)
is a predetermined empirical constant to
convert flow to absolute units of ml/100 g/min
and CBV to absolute units of ml/g. The value of
\( k \), after correction for the difference in field
strength in our study and that of the previous
work from which it was derived, was 1.344 for
the CBF and CBV calculations, and 1.0 for the
MTT calculations. The conversion factor was
extrapolated to 1.5 T by assuming a linear
dependence of the magnitude of susceptibility
contrast upon field strength.¹⁰

DATA ANALYSIS
There were no significant left to right differ-
ences and therefore values from left and right
regions of interest were pooled. Mean grey and
white matter values were obtained by averaging
values from the frontal, parietal, and occipital
cortical regions and the three white matter
regions respectively. Data for the frontal and
occipital grey matter were pooled, as were the
data from the periventricular white matter, and
the same comparisons made as for the
individual ROIs. Mean values were compared
between patients and controls using an un-
paired two tailed student’s t test. In view of the
multiple comparisons made a significance level
of \( p=0.01 \) was used.

Two potential problems which could invali-
date the CBF measurements were blood-brain
barrier breakdown and a ΔR₂ time curve is
delayed relative to the arterial input curve in
some patient subgroups. Therefore data analy-
sis was also performed to determine whether
there was any evidence of white matter blood-
brain barrier leakage in this condition which
would invalidate the assumptions made in CBF
estimation. These assume that the tracer
remains in the intravascular compartment. The
postcontrast–precontrast ΔR₂ difference was
used to indicate leakage. If vascular leakage
does not occur, the signal will return to its pre-
contrast level after the bolus injection, or in the
presence of tracer recirculation, the postbolus
signal will be lower than the prebolus intensity.
If vascular leakage occurs, the postbolus signal
may be increased relative to the prebolus signal.
Once the signal-time curve has been
converted to ΔR₂ values, recirculation would
result in positive ΔR₂, whereas tracer leakage
would then show up as negative postcontrast
ΔR₂.

One possible criticism of the singular value
decomposition technique used to deconvolve
the arterial input curve from the tissue ΔR₂
time curve is its susceptibility to give erroneous
flow values where the ΔR₂ time curve is delayed
relative to the arterial input curve, and if there
is increased dispersion of the contrast agent
bolus before it reaches the tissue.

To investigate these two potential problems
in cases and controls mean tissue signal-times
were calculated for two ROIs placed in the
periventricular white matter of the left and
right hemispheres. The mean arterial input
curve was also calculated. The signal-time
curves were converted into ΔR₂ time curves for
analysis. For each ROI, the mean ΔR₂ was cal-
culated for the precontrast time points, and for
the postcontrast time points. The postcontrast–
precontrast difference in ΔR₂ (diff ΔR₂) was
calculated as an index of vascular leakage. The
mean grey matter and white matter diff ΔR₂
values were then calculated for each subject.
The diff ΔR₂ were compared between the
patients and control group using an unpaired t
test. Tracer arrival time for the AICs and the
tissue ΔR₂ time curves was determined as the
point before ΔR₂ rose above 2×SD of the ΔR₂
of the first 20 time points. The difference in
arrival time (delay) for each region of interest
relative to the AIC was determined, corrected
for delays due to the slice acquisition order of
the multislice echo planar imaging sequence.

Results
Mean CBF measurements are shown in table
1. The controls did not have white matter
abnormalities on T2 weighted images. Typical
images from a control and a patient with leukoaraiosis are shown in figure 1. Mean white matter CBF was significantly reduced in patients by 38% (13.40 (SD 4.87) v 21.74 (SD 3.53) ml/min/100 g, p=0.002). Significant reductions in CBF were seen in all white matter regions; 34% in the anterior, 40% in the superior, and 41% in the posterior white matter. By contrast there was no reduction in mean grey matter CBF. There was no significant reduction in CBF in any grey matter region. There was a wide range of CBF values among the different patients with leukoaraiosis (figure 2). There was no difference in end tidal carbon dioxide concentrations between patients and controls during the acquisition phase of the perfusion imaging (mean (SD) 4.97 (0.51) v 4.97 (0.71) kPa p=0.994).

There was no significant difference in white matter CBV between cases and controls (table 1); mean values were lower in all white matter regions for cases but this did not reach significance for any region. By contrast, mean grey matter CBV was significantly higher in patients than in controls; in each individual region mean values were higher in patients but this only achieved significance for the occipital cortex.

Mean MTT values were higher in all regions of grey and white matter in the patient group, but this only achieved significance for the superior white matter and basal ganglia (table 3).

Table 1 Mean (SD) CBF values (ml/min/100 g) in the two groups

<table>
<thead>
<tr>
<th>Region</th>
<th>Leukoaraiosis (n=8)</th>
<th>Controls (n=9)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean WM</td>
<td>13.40 (4.87)</td>
<td>21.74 (3.53)</td>
<td>0.002</td>
</tr>
<tr>
<td>Mean GM</td>
<td>39.76 (8.38)</td>
<td>40.70 (10.92)</td>
<td>0.85</td>
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<tr>
<td>Mean anterior WM</td>
<td>13.65 (5.93)</td>
<td>20.74 (3.42)</td>
<td>0.01</td>
</tr>
<tr>
<td>Mean posterior WM</td>
<td>11.79 (4.67)</td>
<td>19.81 (4.61)</td>
<td>0.003</td>
</tr>
<tr>
<td>Mean superior WM</td>
<td>14.75 (4.64)</td>
<td>24.66 (4.24)</td>
<td>0.000</td>
</tr>
<tr>
<td>Mean frontal GM</td>
<td>41.85 (10.67)</td>
<td>45.59 (12.23)</td>
<td>0.51</td>
</tr>
<tr>
<td>Mean parietal GM</td>
<td>40.63 (8.63)</td>
<td>42.12 (11.78)</td>
<td>0.77</td>
</tr>
<tr>
<td>Mean occipital GM</td>
<td>36.51 (7.08)</td>
<td>34.38 (9.47)</td>
<td>0.55</td>
</tr>
<tr>
<td>Mean basal ganglia</td>
<td>44.57 (11.12)</td>
<td>52.11 (16.40)</td>
<td>0.28</td>
</tr>
</tbody>
</table>

WM=White matter; GM=grey matter. Significant differences are highlighted in bold. A significance level of 0.01 has been used in view of the multiple comparisons made.

Figure 2 Mean white matter CBF values in controls and patients with ischaemic leukoaraiosis.
Table 3 Mean (SD) MTT values (s) in the two groups

<table>
<thead>
<tr>
<th>Region</th>
<th>Leukoaraiosis (n=8)</th>
<th>Controls (n=9)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean WM</td>
<td>1.74 (0.48)</td>
<td>2.03 (0.44)</td>
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<tr>
<td>Mean GM</td>
<td>3.92 (0.93)</td>
<td>3.07 (0.63)</td>
<td>0.01</td>
</tr>
<tr>
<td>Mean anterior WM</td>
<td>1.83 (0.50)</td>
<td>2.14 (0.48)</td>
<td>0.21</td>
</tr>
<tr>
<td>Mean posterior WM</td>
<td>1.69 (0.54)</td>
<td>1.99 (0.47)</td>
<td>0.24</td>
</tr>
<tr>
<td>Mean superior WM</td>
<td>1.71 (0.47)</td>
<td>1.97 (0.41)</td>
<td>0.26</td>
</tr>
<tr>
<td>Mean frontal GM</td>
<td>3.88 (1.13)</td>
<td>3.18 (0.67)</td>
<td>0.16</td>
</tr>
<tr>
<td>Mean parietal GM</td>
<td>3.92 (0.92)</td>
<td>3.11 (0.64)</td>
<td>0.06</td>
</tr>
<tr>
<td>Mean occipital GM</td>
<td>3.96 (0.84)</td>
<td>2.92 (0.49)</td>
<td>0.01</td>
</tr>
<tr>
<td>Mean basal ganglia</td>
<td>4.31 (1.61)</td>
<td>3.36 (0.91)</td>
<td>0.17</td>
</tr>
</tbody>
</table>

Explanations as in table 1.

There was no difference in white matter diffusivity (AR, values in patients versus controls (0.010 (SD 0.07)) vs. 0.08 (SD 0.01)) (p=0.7). There was no significant delay in tracer arrival time in the white matter of patients compared with controls (2.77 (SD 1.34) vs. 2.83 (SD 1.63)) s, p=0.94.

Discussion

Using a quantitative MR perfusion method we have shown reduced white matter CBF in patients with ischaemic leukoaraiosis, but normal grey matter CBF. The absolute values of CBF we obtained in the patient groups were very similar to those in previous PET studies in which both white and grey matter flow have been determined. This provides further evidence for the validity of the rCBF measurements using this quantitative MR perfusion technique.

We found no significant difference in white matter CBV between the two groups although mean values were non-significantly lower for all white matter regions in the leukoaraiosis group. However, our study was not sufficiently powered to determine if this is a real difference. Power calculations derived using the SD from the control group in this study showed that a sample size of about 100 would be needed to show a significant difference of this magnitude. The results we obtained were very similar to those in a previous PET study in which patients with white matter leukoaraiosis (n=8) were obtained. In our study we derived this from the middle cerebral artery; this was possible in all cases. An alternative is to derive it from the internal carotid artery but this was not possible in our images due to susceptibility artefact in this region. We used an SE sequence as this has better sensitivity and selectivity for CBF in the microvasculature. With gradient echo EPI, the dependence of the rCBF measurements on vessel radius increases, and is considerably larger for arteries than for small arterioles. In SE-EPI, CBF reaches a peak for vessel radius of less than 10 μm in arteries and then falls off with increasing vessel size. In addition the position of the peak CBF moves to a smaller vessel radius with increasing contrast agent dose. Therefore the CBF signal derived from SE-EPI is predominantly from the microvasculature whereas that from gradient echo EPI is also derived to a greater extent from the larger vessels.

Estimation of cerebral perfusion using dynamic susceptibility imaging assumes that the contrast agent remains in the intravascular compartment. If blood-brain barrier disruption and vascular leakage occurred this would invalidate the measurements. Blood-brain barrier disruption has been reported in patients with leukoaraiosis who have been reported to have higher CSF/serum albumin ratios, and this would be consistent with blood-brain barrier breakdown. If vascular leakage does not occur, the signal will return to its precontrast level after the bolus injection, or in the presence of tracer recirculation, the postbolus signal will be lower than the prebolus intensity. We found no difference in the postcontrast-to-precontrast AR difference between the patient and control group, suggesting leakage does not occur at least during the time scale of the perfusion measurement. This is consistent with a recent study in which patients with white matter...
vascular lesions, some of whom had functional blood-brain barrier disruption as evidenced by increased CSF/serum albumin ratios, were imaged repeatedly over a 30 minute period just before and after a double standard dose of a paramagnetic contrast medium, using a T1 weighted sequence. There was no significant change in the MR signal in the white matter lesions during this period. Another possible criticism of the singular value decomposition technique used to deconvolve the arterial input curve from the tissue AR, time curve is its susceptibility to give erroneous flow values where the AR, time curve is delayed relative to the arterial input curve. However, we found no difference in the mean delay between patients and controls in the white matter.

In summary, quantitative exogenous perfusion MRI allows absolute quantification of perfusion in diffuse cerebrovascular disease such as ischaemic leuкоaraiosis with a resolution similar to or better than previous methods. Using this technique our results provide further evidence for the role of chronic ischaemia in this condition.

This work was supported by a grant from the Stroke Association of the United Kingdom. We thank the neuroimaging staff and in particular Amanda Glover and Caroline Andrews for their assistance in acquiring the MR data. We thank Professor Steve Jackson for allowing us to study the health care of the elderly normal control database.