Do white matter changes on MRI and CT differentiate vascular dementia from Alzheimer’s disease?

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SUMMARY MRI showed white matter changes in all 29 patients with vascular dementia and in eight out of 22 patients with Alzheimer’s disease. The corresponding figures for CT were 26 and 1, respectively. White matter changes are therefore a useful diagnostic aid in the differential diagnosis between vascular dementia and Alzheimer’s disease.

Vascular dementia and Alzheimer’s disease are the two main types of dementia.¹ Neuroradiological differentiation of vascular dementia and Alzheimer’s disease has usually focused on differences in cortical and central atrophic changes² ³ and on the presence or absence of infarcts on CT.³–⁶ CT, however, has not differentiated the types of dementia adequately.²–⁷ obviously because white matter changes have often been neglected. Our preliminary experience with MRI showed white matter changes in vascular dementia, but not in Alzheimer’s disease.⁸ A larger series of consecutive patients with vascular dementia or Alzheimer’s disease was studied by MRI and CT in order to evaluate white matter changes in these types of dementia and the possible usefulness of MRI as their differential diagnostic aid.

Patients and methods

In the Memory and Dementia Project at the Department of Neurology, University of Helsinki, 233 consecutively admitted patients with dementia were studied from 4 May, 1982 to 31 May, 1985. Alzheimer’s disease comprised 68 patients, vascular dementia 133 and secondary causes of dementia 32 patients. From this source population 51 patients, fulfilling the criteria of vascular dementia or Alzheimer’s disease, and consecutively studied by both MRI and CT were included in the present study. There were 29 patients with vascular dementia (10 men, 19 women) and 22 patients with Alzheimer’s disease (12 men, 10 women). The mean age (mean ± SEM) was 68.7 ± 1.7 years (range 53 to 87) in patients with vascular dementia and 64.0 ± 1.7 years (range 50 to 77) in those with Alzheimer’s disease.

All patients were clinically examined by two neurologists (TE, RS). Diagnostic assessment included detailed medical history, physical and neurological investigation, screening laboratory tests, investigation of the cerebrospinal fluid, neuropsychological test battery, EEG and CT.

The clinical criteria for vascular dementia were: dementia with a history of focal or acute signs or symptoms referable to disturbances in cerebral blood circulation, often with elevated blood pressure, abrupt onset of symptoms, stepwise deterioration and relative preservation of personality.⁹ ¹⁰ The corresponding criteria for Alzheimer’s disease were: dementia with insidious onset and progressive deteriorating course without history, signs or symptoms of focal brain damage and without history or laboratory findings referable to any other aetiology.¹¹ Secondary causes of dementia, other than vascular aetiology, including normal pressure hydrocephalus,¹² ¹³ and factors known to give reduction in white matter attenuation on CT, for example, multiple sclerosis and leukodystrophies,¹⁴–¹⁷ were excluded. The CT data together with clinical data were used in the exclusion of secondary causes of dementia.

Dementia was assessed using the Dementia Scale¹⁸ and a Luria based neuropsychological test battery¹⁹ according to the DSM III criteria.⁹ Dementia was mild in 10 patients with vascular dementia and in 10 patients with Alzheimer’s disease, moderate in 13 with vascular dementia and in nine with Alzheimer’s disease and severe in six with vascular dementia and in three with Alzheimer’s disease. The mean duration (± SEM) of cognitive symptoms was 3.3 ± 0.5 years (range 0 to 11) in vascular dementia and 4.0 ± 0.5 years (range 1 to 9) in Alzheimer’s disease.

The Modified Ischemic Stroke Score²⁰ was also given to the patients. It was 7.3 ± 0.5 (mean ± SEM, range 2 to 12) in patients with vascular dementia and 0.3 ± 0.1 (range 0 to 1) in those with Alzheimer’s disease (p < 0.001).
MRI was performed using a prototype MRI equipment manufactured by Instrumentarium Corp., Helsinki, Finland. The device operated with a main magnetic field of 0.17 T (corresponding to 7.13 MHz) produced by a superconducting magnet. Axial transverse sections approximately 10 mm thick were obtained and displayed on a matrix of 128 x 128 pixels. Data acquisition time varied from 3.5 to 9 min. To obtain T1-weighted images, an IR 1500(400)/30 sequence was used and for T2-weighted images, the SE 2000/100 sequence was used. Noncontrast CT was performed with an EMI CT 1010 or Somatom DR2 scanners. The time interval between MRI and CT was less than 3 months and during this time no clinical acute events were permitted.

In the T2 weighted images white matter changes and areas of increased signal intensity compatible with infarcts were recorded independently by two of us (LK, JS) without knowledge of the clinical diagnosis. Areas of increased signal intensity relative to the rest of the brain, which are presumed to be due to longer T2, around frontal or occipital horns, or along the bodies of the lateral ventricles were recorded. The lesion was rated to be mild if there was any sign of long T2 in the white matter extending less than one fourth of the total white matter area (fig, a). Changes were rated moderate if long T2 was up to half of the white matter area (fig, b), and severe if the long T2 extended over one half of the total white matter area (fig, c). On noncontrast CT images the white matter low attenuations and infarcts were rated by two neuroradiologists (LK, MV) using the same criteria (fig, d–f). Statistical differences between the groups were calculated using the two-tailed t test and chi-square test. The predictive values of diagnostic test were also calculated.21

Fig. T2 weighted MR image demonstrating mild (a) and moderate (b) degree of white matter change around the frontal and occipital horns. The lesion appears as high signal intensity (arrows). Plain CT scans (d, e) with low attenuation in the corresponding areas. T2 weighted MR image (c) of a patient with severe white matter change appearing as high signal intensity in the periventricular white matter. Corresponding CT image (f) demonstrates extensive low attenuation in the same area.
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Table 1. White matter (WM) changes in vascular dementia (VD) (n = 29) and Alzheimer's disease (AD) (n = 22) by MRI and CT in different brain regions.

<table>
<thead>
<tr>
<th>Grade of WM changes</th>
<th>Frontal</th>
<th>Periventricular</th>
<th>Occipital</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>MRI VD</td>
<td>AD CT</td>
<td>MRI VD</td>
</tr>
<tr>
<td>None</td>
<td>0</td>
<td>17</td>
<td>3</td>
</tr>
<tr>
<td>Mild</td>
<td>7</td>
<td>4</td>
<td>10</td>
</tr>
<tr>
<td>Moderate</td>
<td>8</td>
<td>1</td>
<td>8</td>
</tr>
<tr>
<td>Severe</td>
<td>5</td>
<td>1</td>
<td>10</td>
</tr>
</tbody>
</table>

Table 2. Cerebral infarcts and white matter (WM) changes in vascular dementia (VD) and Alzheimer's disease (AD) by MRI and CT.

<table>
<thead>
<tr>
<th>Infarcts</th>
<th>WM changes</th>
<th>Infarcts or WM changes</th>
</tr>
</thead>
<tbody>
<tr>
<td>MRI VD</td>
<td>AD CT</td>
<td>MRI VD</td>
</tr>
<tr>
<td>No</td>
<td>10</td>
<td>22</td>
</tr>
<tr>
<td>Yes</td>
<td>19</td>
<td>0</td>
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</table>

Table 3. Sensitivity, specificity, positive and negative predictive values (%) of white matter (WM) changes on MRI and CT in the diagnosis of vascular dementia.

<table>
<thead>
<tr>
<th>Frontal WM</th>
<th>Periventricular WM</th>
<th>Occipital WM</th>
</tr>
</thead>
<tbody>
<tr>
<td>MRI VD</td>
<td>100.0</td>
<td>89.7</td>
</tr>
<tr>
<td>CT</td>
<td>96.6</td>
<td>51.7</td>
</tr>
<tr>
<td>Sensitivity</td>
<td>96.6</td>
<td>51.7</td>
</tr>
<tr>
<td>Specificity</td>
<td>96.6</td>
<td>51.7</td>
</tr>
<tr>
<td>Positive predictive value</td>
<td>96.6</td>
<td>51.7</td>
</tr>
<tr>
<td>Negative predictive value</td>
<td>96.6</td>
<td>51.7</td>
</tr>
</tbody>
</table>

*Computed from data given in Table 1.

Table 4. Sensitivity, specificity, positive predictive values (%) and negative predictive values (%) of infarcts and white matter (WM) changes on MRI and CT in the diagnosis of vascular dementia.

<table>
<thead>
<tr>
<th>Infarcts</th>
<th>WM changes</th>
<th>Infarcts or WM changes</th>
</tr>
</thead>
<tbody>
<tr>
<td>MRI VD</td>
<td>AD CT</td>
<td>MRI VD</td>
</tr>
<tr>
<td>Sensitivity</td>
<td>100.0</td>
<td>89.7</td>
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<tr>
<td>Specificity</td>
<td>100.0</td>
<td>89.7</td>
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<tr>
<td>Positive predictive value</td>
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<td>89.7</td>
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<tr>
<td>Negative predictive value</td>
<td>100.0</td>
<td>89.7</td>
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*Computed from data given in Table 2.

Results

White matter changes on MRI and CT are shown in Table 1. On MRI white matter changes were detected in all patients with vascular dementia, but only in eight patients with Alzheimer's disease. The degree of the white matter changes was moderate to severe in vascular dementia, while it was mostly mild in Alzheimer's disease. MRI was more sensitive to detect white matter changes than CT.

White matter changes and areas of altered signal compatible with cerebral infarcts as seen on MRI and CT are shown in Table 2. Nineteen (65.5%) of the patients with vascular dementia had infarcts on MRI and 18 (62.0%) on CT, while none of the patients with Alzheimer's disease had infarcts on MRI or CT. Infarcts or white matter changes were seen in all of the 29 patients with vascular dementia on MRI and in 28 on CT. One patient with vascular dementia who had no vascular changes on CT at the time of in-
vestigation developed an infarct visible on CT three months later.

The presence or absence of white matter changes and infarcts on MRI and CT were further evaluated by calculating sensitivity, specificity, positive and negative predictive values of these changes in the diagnosis of vascular dementia (tables 3 and 4). Frontal white matter changes were most sensitive, while occipital changes were most specific. If mild white matter changes were used as a positive indicator of vascular dementia, MRI gave high sensitivity, but lower specificity. On the other hand, CT gave better specificity but lower sensitivity. If only moderate to severe white matter changes were regarded to be indicative of vascular dementia, the specificity improved at the cost of sensitivity. Infarcts gave better specificity but lower sensitivity than white matter changes alone or combined with infarcts. Either white matter changes or infarcts as an indicator of vascular dementia on CT gave good sensitivity (97%) and specificity (96%). On MRI these changes, however, did not give a better discriminative power than the white matter changes alone (sensitivity 100%, specificity 63%).

White matter changes on MRI and CT by age and severity of dementia are given in table 5. The number of patients with white matter changes increases with age (MRI: \( \chi^2 = 8.36, p = 0.015 \); CT: \( \chi^2 = 6.22, p = 0.045 \)), but not with the severity of dementia.

**Discussion**

All the 29 patients with vascular dementia had white matter changes on MRI, while 26 of them showed changes on CT. Eight of the 22 patients with Alzheimer's disease had white matter changes on MRI, and only one on CT. White matter changes in Alzheimer's disease were mostly mild. The eight Alzheimer patients with white matter changes were older and had dementia of mild to moderate degree. Compared to CT, MRI was more sensitive to reveal white matter changes both in vascular dementia and Alzheimer's disease. It especially gave a good contrast on the T2-weighted MRI, as noted also previously.8 17 22 The overall correlation between MRI and CT was good. Detection of white matter changes is better with the high resolution CT scanner, but comparisons between the two CT scanners used in the present study were not made in detail. The same is true with the MR imagers with thinner sections and better spatial resolution.

None of the Alzheimer patients showed infarcts on MRI or CT. Of the patients with vascular dementia 19 (65.5%) had infarcts on MRI and 18 (62.0%) on CT. These figures are higher than in previous studies where the corresponding figures on CT varied from 26% to 42%.3–6 The appearance of white matter changes or infarcts on MRI and on CT in the present series suggests that MRI is more sensitive than CT to demonstrate these changes.

White matter changes have been described previously in disorders related to different types of vascular dementia includingBinswanger's disease, the lacunar state, systemic hypoperfusion and multiple strokes.8 17 23–27 In this study all patients with vascular dementia had white matter changes. The loss of total brain volume due to infarcts in vascular dementia has been emphasised in the development of the dementia syndrome.28 However, our present experience suggests that also disconnection to the cortex due to white matter changes may give rise to symptoms of vascular dementia.

In our preliminary report of three presenile Alzheimer patients we did not find any white matter changes either on CT or on MRI,8 but in the present larger series we found mild to moderate white matter changes in some Alzheimer patients. Loss of greywhite matter discriminability29 and decreased white matter parenchymal density, with no clear white matter low attenuation, have been reported on CT of patients with Alzheimer's disease.30 Decreased white matter density, however, was not confirmed by Gado et al.31 Previously MRI was used in the differentiation between vascular dementia and Alz-
heimer’s disease. In that study proton density values in white matter were lower in vascular dementia than in Alzheimer’s disease while T1 values were equally lowered in both types of dementia. However, T2 values, which were more informative in our present study, were not used. Therefore, our results are not comparable with those of Besson et al.

Some white matter changes might be age related. Zatz et al reported decreased attenuation values of the white matter on CT in normal elderly people. However, no clear focal or generalised low attenuation of white matter was noted. Young et al pointed out that small areas of long T2 around the frontal horns might be normal. In our series the occurrence of white matter changes increased with age but not with the severity of dementia. Thus the mild white matter changes among Alzheimer patients may be partly age related. White matter changes among the patients with mild to moderate Alzheimer's disease may also be related to minor ischaemic events. A further possibility is that these older Alzheimer patients with white matter changes on MRI constitute its own diagnostic entity.

In conclusion, white matter changes as seen on MRI and also on CT are a useful aid in the differential diagnosis between the two main types of dementia: vascular dementia and Alzheimer’s disease.

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