Hippocampal MRI volumetry in cognitively discordant monozygotic twin pairs

T Järvenpää, M P Laakso, R Rossi, M Koskenvuo, J Kaprio, I Räihä, T Kurki, M Laine, G B Frisoni, J O Rinne

Objective: To investigate whether hippocampal atrophy, a proxy for incipient Alzheimer’s disease, can be detected in non-demented monozygotic co-twins of demented twins by using volumetric magnetic resonance imaging (MRI).

Methods: Seven pairs of monozygotic female twins discordant for cognitive function (mean [SD] age 75 (4) years), and 10 age and education matched healthy controls (seven women, three men; mean age 73 (3) years) were studied with volumetric MRI.

Results: The mean normalised right hippocampal volume was 31% lower (p = 0.002) in the demented twins, and 6% lower (p = 0.45) in the non-demented twins than in the controls. In the left hippocampus, the mean normalised volume was 36% lower (p<0.001) in the demented twins, and 9% lower (p = 0.13) in the non-demented twins than in the controls.

Conclusions: Significant hippocampal atrophy was detected in the demented twins compared with the controls. This is in line with previous imaging and pathological studies, with hippocampus showing the early changes in Alzheimer’s disease. In the non-demented twins, only a minor, non-significant reduction was observed in the hippocampal volumes compared with the controls. This could reflect gene–environment interactions that have protected the non-demented twins longer than their demented co-twins and contributed to the relative preservation of their hippocampal volumes, or it could be a sign of preclinical Alzheimer’s disease in the non-demented twins.
right handed. The mean (SD) age of the twins was 75 (4) years. The mean mini-mental state examination (MMSE) score of the demented twins was 21.1 (6.0) and that of their non-demented co-twins, 26.0 (2.8). All twins were screened for hypothyroidism and vitamin B-12 deficiency to exclude these conditions as possible causes of dementia. No clinically significant abnormalities that could have affected cognitive function in the twins were found. None of the twins had experienced any serious head trauma, had a history of epilepsy, or had experienced single seizures or episodes of status epilepticus. This was confirmed from the medical records. In the interview with the twins and informants, no history of prolonged febrile convulsions in childhood became apparent. Furthermore, no major depression was detected in any twins, based on the judgement of the clinician or neuropsychologist; possible minor depressive symptoms were not likely to have confounded the neuropsychological test results.

Confirmation of zygosity and ApoE status
The zygosity was confirmed by DNA extracted from venous blood samples. Ten highly polymorphic genetic markers used by the paternity testing laboratory of the National Public Health Institute were determined, and monozygosity was inferred if no intrapair differences in the markers were observed. Three twin pairs had an ApoEε4 allele, with one twin being homzygous for this allele. The remaining four pairs were homzygous for the ε3 allele.

Neuropsychological testing
In addition to the initial TELE interview, the twins’ cognitive discordance was confirmed with a comprehensive neuropsychological test battery including the Wechsler memory scale–revised (logical memory immediate and delayed, visual reproduction immediate and delayed), Wechsler memory scale (digit span), Wechsler adult intelligence scale–revised (similarities, block design), verbal fluency (animals and letter s), trail making A, and Boston naming test. These tests assess memory, attention, language, psychomotor functions, and visuospatial and visuoconstructive abilities. The results of the above mentioned tests in 113 neurologically healthy elderly Finnish individuals of comparable age were used as reference values. If the individual had a subjective memory complaint, scored at least 1.5 SD under the mean reference value in any cognitive domain, had normal general cognitive function, and had preservation of the activities of daily living, the criteria for mild cognitive impairment were fulfilled. If the individual showed widespread impairment (scored at least 1.5 SD under the mean reference value in any other cognitive domain in addition to the memory deficit), and had a progressive decline in memory and other cognitive functions over time, the NINCDS-ADRSA criteria for probable Alzheimer’s disease were fulfilled. All of the seven twin pairs were cognitively intact. According to the above mentioned criteria, in four pairs one co-twin was cognitively intact and the other was demented. In two pairs one co-twin was intact, and the other had mild cognitive impairment. In the remaining pair, one co-twin had mild cognitive impairment, and the other was demented. In addition to the above mentioned tests, the twins underwent the Consortium to Establish a Registry for Alzheimer’s Disease (CERAD) tests.

Controls
The control group consisted of 10 healthy volunteers (seven women and three men) with no previous history of neurological or psychiatric diseases. These controls were age and education matched. The duration of education was categorised into four classes (<6 years, 7–9 years, 10–12 years, and >12 years). There was no difference in the distribution of education between the demented twins and the controls (p = 1.00, Fisher’s exact test), between the non-demented twins and controls (p = 1.00, Fisher’s exact test), or between the demented and non-demented twins (p = 0.32, McNemar’s test). The controls underwent the same neuropsychological examination as the twins. The neuropsychological test scores of the control group were also compared with the reference values of the 113 neurologically healthy elderly Finnish individuals, and after this thorough examination the controls were confirmed as cognitively intact. All controls were right handed. The mean age of the control group was 73 (3) years, and their mean MMSE score was 28.5 (1.1).

Written informed consent was obtained from all participants before the MRI and neuropsychological investigation. The study protocol was approved by the joint ethics committee of the University of Turku and the Turku University Central Hospital.

MRI
MRI scanning
The participants were scanned with a 1.5 Tesla MRI device (three controls with a Siemens Magnetom, Erlangen, Germany; seven controls and all twins with GE Signa, General Electric Medical Systems, Milwaukee, Wisconsin, USA) at the Turku University Central Hospital. For the Siemens’ scanner, we used a three dimensional, magnetisation prepared, rapid acquisition gradient echo sequence (time of repetition (TR) 10 ms, time of echo (TE) 4 ms, flip angle 10°, matrix 256×256, one acquisition), which resulted in 128 1.5 mm thick sagittal images with no interslice gap. For GE Signa, we used three dimensional, fast spoiled, gradient echo sequence (TR 11.3 ms, TE 4.2 ms, flip angle 20°, matrix 256×256, one acquisition), which resulted in 124 1.2 mm thick axial images with no interslice gap.

To measure the hippocampal volumes, the scans were saved on an optical disc and analysed at the Kuopio University Hospital (Kuopio, Finland). First, the scans were reconstructed into 2.0 mm thick contiguous coronal slices oriented perpendicularly to the intercommissural line. The hippocampi were manually traced slice by slice by one person, blinded to the subjects’ clinical data, using custom made software for a standard Siemens’ work console. The rostral end of the hippocampus, when it first appears below the amygdala, was the anatomical starting point. Tracing of the hippocampus ended posteriorly in the section where the crura of the fornices depart from the lateral wall of the lateral ventricles.

The intraclass correlation coefficient for intrarater reliability was 0.95. The intracranial area on a coronal section at the level of the anterior commissure was measured and used for normalisation of the volumetric data. For data presentation purposes, the volumes reported herein are normalised to the intracranial area according to the formula: [volume/intracranial area]×1000.

Statistical analyses
The results are shown as mean (SD). The equality in the distribution of education between the demented twins and controls, and between the non-demented twins and controls, was tested with Fisher’s exact test. McNemar’s test was used in the comparison of education between the demented and non-demented twins, as the twins were considered to be correlated (paired) samples. Comparisons between the hippocampal volumes of the demented and non-demented co-twins were done with paired t tests. Differences in the hippocampal volumes between the demented twins and controls, and between the non-demented twins and controls, were evaluated with two sample t tests. The associations between the neuropsychological test scores and hippocampal...
Volumes were analysed with Pearson’s correlation coefficients. In all tests, differences were considered significant at a probability (p) value of <0.05. Statistical analyses were done with the SAS System for Windows, release 8.02 (SAS Institute, Cary, North Carolina, USA).

### RESULTS

The mean normalised right and left hippocampal volumes in the demented twins, their non-demented co-twins, and the controls are shown in Table 1. In the right hippocampus, the mean hippocampal volume was 31% lower in the demented twins (t = −3.66, df = 15, p = 0.002), and 6% lower in the non-demented twins (t = −0.77, df = 15, p = 0.45) than in the controls. When the demented twins were compared with their non-demented co-twins, a significant 27% reduction (t = −5.10, df = 6, p = 0.002) was found in the mean right hippocampal volume. In the left hippocampus, the mean hippocampal volume was 36% lower in the demented twins (t = −5.79, df = 15, p < 0.001), and 9% lower in the non-demented twins (t = −1.59, df = 15, p = 0.13) than in the controls. Here, a significant 29% reduction (t = −6.17, df = 6, p < 0.001) was also found in the demented twins compared with their non-demented co-twins. Figure 1 shows the scatterplots of normalised right and left hippocampal volumes in the three groups. Correlation analysis revealed no significant associations between the hippocampal volumes and MMSE scores, or delayed recall and savings scores of the demented twins, their non-demented co-twins, and the controls. This is in accordance with previous twin studies, in which MRI findings were either accompanied by cognitive impairment, or else no pathological MRI findings were found despite cognitive impairment in the unaffected co-twins. However, in only one study was medial temporal lobe atrophy measured quantitatively, while the other two studies concentrated on visual assessment of cortical atrophy without any focus on the medial temporal lobe. Thus the earliest changes of medial temporal lobe atrophy, before the first signs of cognitive deterioration, might have been missed.

In our study, significant hippocampal atrophy was detected in the demented twins compared with their non-demented co-twins and controls. This is in accordance with previous studies in which MRI volumetry has been shown to be a sensitive means of detecting hippocampal atrophy early in the disease process of Alzheimer’s disease, and to differentiate patients with Alzheimer’s disease from healthy controls. In the non-demented twins in our study only a minor 6–9% reduction in hippocampal volumes was observed compared with the controls. Earlier studies have shown that volumetric MRI measurements of the hippocampus have been 11–15% lower in subjects with mild cognitive impairment than in healthy controls. However, all but one of the non-demented twins in our study were diagnosed as being cognitively intact. Thus this is probably the reason why we did not find any significant volume losses in our study, in contrast with those involving subjects with mild cognitive impairment.

We did not find any significant differences in mean hippocampal volumes between the non-demented twins and the controls, but issues that should be considered are the relatively small number of twin pairs and the large within-group variation in hippocampal volumes. It is not likely that the two scanners used in this study contributed to the within-group variation, given that all the twin pairs were scanned with the same scanner. In one previous study, co-registered serial MRI showed an accelerated atrophy rate, in comparison with controls, in the medial temporal lobe of presymptomatic subjects who carried a known autosomal dominant mutation leading to early onset Alzheimer’s disease. Thus a longitudinal follow up would be needed to show whether the non-demented twins in our study would have a greater atrophy rate than the controls, or whether those who might convert to Alzheimer’s disease would have even greater hippocampal atrophy. In our study, the correlations between the cognitive test scores and hippocampal volumes in the demented twins were positive but did not reach statistical significance, unlike in some previous studies on patients with Alzheimer’s disease, which probably

### DISCUSSION

This is the first volumetric MRI study, to our knowledge, that has been carried out in a group of monozygotic twins discordant for cognitive function. We found the mean normalised right and left hippocampal volumes of the demented twins to be significantly reduced, in a range of 27% to 36% compared with their non-demented co-twins and controls. The non-demented twins showed a 6–9% hippocampal volume loss compared with the controls, which did not reach statistical significance.

Previously, three brain MRI studies have been done on monozygotic twins discordant for Alzheimer’s disease, and each study has been a case report based on a single pair. Thus no statistical conclusions could have been drawn from those studies. In a report by Luxenberg et al in one male monozygotic twin pair discordant for Alzheimer’s disease, MRI of the affected twin showed marked loss of cortical grey matter, enlargement of sulcal spaces, and ventricular dilatation, while MRI in the unaffected co-twin was normal. In neuropsychological tests the affected twin had marked deficits in all areas of cognition, while the unaffected co-twin had milder impairment. Small et al studied one female monozygotic twin pair appearing discordant for Alzheimer’s disease. MRI showed atrophy in both twins, with larger amounts of sulcal widening, ventricular enlargement, and loss of normal contour of the lateral ventricles in the affected twin. In neuropsychological tests the affected twin showed extensive deterioration, while her unaffected co-twin had milder cognitive decline. In a study by Geroldi and associates with one female monozygotic twin pair discordant for Alzheimer’s disease, the MRI showed medial temporal lobe atrophy only in the affected twin. Cognitive functions were normal in the unaffected co-twin. Thus in these previous twin studies, pathological MRI findings were either accompanied by cognitive impairment, or else no pathological MRI findings were found despite cognitive impairment in the unaffected co-twins. However, in only one study was medial temporal lobe atrophy measured quantitatively, while the other two studies concentrated on visual assessment of cortical atrophy without any focus on the medial temporal lobe. Thus the earliest changes of medial temporal lobe atrophy, before the first signs of cognitive deterioration, might have been missed.

### Table 1: Normalised right and left hippocampal volumes in the demented twins, their non-demented co-twins, and the controls

<table>
<thead>
<tr>
<th></th>
<th>Demented twins (n=7)</th>
<th>Non-demented co-twins (n=7)</th>
<th>Controls (n=10)</th>
<th>p Values</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Volume (SD)</td>
<td>%</td>
<td>Volume (SD)</td>
<td>%</td>
</tr>
<tr>
<td>Right</td>
<td>102.4 (23.8)</td>
<td>69</td>
<td>139.4 (14.6)</td>
<td>94</td>
</tr>
<tr>
<td>Left</td>
<td>90.3 (19.0)</td>
<td>64</td>
<td>127.8 (15.9)</td>
<td>91</td>
</tr>
</tbody>
</table>

Values are mean (SD). Percentages indicate the proportion of the control mean value.

1Statistical method: paired t test
2Statistical method: two sample t test

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The scatterplots of normalised volumes (hippocampal volume/intracranial area) in the right and left hippocampus in the demented twins, their non-demented co-twins, and the controls. Each twin pair is connected with a line. The horizontal line shows the mean normalised hippocampal volume in each group.

REFERENCES

Reference linking to full text of more than 200 journals
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