A randomised placebo controlled study to assess the effects of cholinergic treatment on muscarinic receptors in Alzheimer’s disease


**Objective:** To determine the effects of cholinergic treatment on the muscarinic receptor in patients with Alzheimer’s disease.

**Methods:** 12 patients with mild to moderate Alzheimer’s disease and six controls were studied. The patients underwent ADAS-COG psychometric assessment and SPECT brain imaging with $^{123}$I quinuclidinyl benzilate (QNB), to demonstrate the postsynaptic muscarinic M1 receptor, before being randomised in a double blind study to receive either an acetylcholinesterase inhibitor (donepezil) or placebo for four months. Following this, the ADAS-COG and the $^{123}$I-QNB receptor scan were repeated. The controls were imaged on one occasion only. All image analyses were undertaken using SPM99.

**Results:** $^{123}$I-QNB imaging showed a significant relation between baseline psychometric impairment and deficits on scanning. Both placebo and actively treated groups had reductions in $^{123}$I-QNB uptake. Greater reductions in receptor binding were demonstrated in the placebo group than in those receiving active treatment. Intraindividual reproducibility of the $^{123}$I-QNB imaging technique appeared highly robust.

**Conclusions:** The results suggest that $^{123}$I-QNB uptake is better preserved in Alzheimer’s disease patients on cholinergic treatment than on placebo. Cholinergic treatment may play a neuroprotective role. Sequential $^{123}$I-QNB imaging seems to be a powerful tool in monitoring the response of these receptors to disease modifying treatments.

**METHODS**

**Recruitment**

Twelve patients who fulfilled the NINCDS-ARDRA criteria for probable Alzheimer’s dementia were recruited for this study at the memory clinic, Moorgreen Hospital, Southampton, by a consultant in old age psychiatry. Six healthy controls without a history of head injury or neuropsychiatric illness, and with no apparent cognitive impairment, were also recruited.

**Trial design**

The 12 patients with Alzheimer’s disease underwent baseline ADAS-COG (Alzheimer’s disease assessment scale—cognitive behaviour) psychometric assessment and $^{123}$I-QNB imaging followed by randomisation in a double blind trial to receive either four months of treatment with an AChE-I (donepezil 5 mg daily increasing to 10 mg at four weeks) or placebo. Following this period, and before discontinuation of the allocated treatment, the psychometric assessment and brain scan were repeated. The six controls were imaged on one occasion only and did not undergo formal psychometric testing.

**$^{123}$I-QNB preparation and imaging**

The (R,R) QNB isomer was synthesised by the Department of Radiopharmacy, Glasgow University, and was subsequently labelled locally with $^{123}$I using a high performance liquid chromatographic technique. Five hours after the intravenous administration of 160 MBq $^{123}$I-QNB, the subjects underwent a 30 minute tomographic acquisition on a SMV DST-XL dual head gamma camera. These projections were prefiltered, corrected for decay and attenuation, and reconstructed with a ramp filter. The statistical parametric software package (SPM99) was used for image analysis. The reconstructed images were registered to a single photon emission computed tomographic (SPECT) template image set in standardised stereotactic space, smoothed, and normalised to the mean count within the image.

**Image analyses**

The following analyses of the SPM maps of the $^{123}$I-QNB images were undertaken:

- Assessment of associations between the $^{123}$I-QNB images and the psychometric assessment scores using linear regression analysis;
- Group comparison ($t$ test) of the baseline patient images vs the controls;
- Group comparison ($t$ test) of the baseline images of the actively treated patient group vs the placebo group;
RESULTS
The median ages of the patient and control groups were 75 years (range 58 to 87) and 70 years (range 65 to 79), respectively; there was no significant difference between medians (Mann-Whitney test, p = 0.20). The ADAS-COG scores of two of the patients in this pilot study were excluded, one because of visual impairment and one because of withdrawal from the trial. The median ADAS-COG scores at baseline of the actively treated and placebo group were 23 (range 11 to 34) and 15 (range 9 to 28), respectively; there was no significant difference between medians (Mann-Whitney test, p = 0.05). The ADAS-COG scores of two of the patients in this pilot study were excluded, one because of visual impairment and one because of withdrawal from the trial. The median ADAS-COG scores at baseline of the actively treated and placebo group were 22 (range 11 to 34) and 15 (range 9 to 28), respectively; there was no significant difference between medians (Mann-Whitney test, p = 0.05). The ADAS-COG scores of two of the patients in this pilot study were excluded, one because of visual impairment and one because of withdrawal from the trial. The median ADAS-COG scores at baseline of the actively treated and placebo group were 23 (range 11 to 34) and 15 (range 9 to 28), respectively; there was no significant difference between medians (Mann-Whitney test, p = 0.27).

All 12 patients successfully underwent the baseline brain imaging and were randomised to receive either a cholinesterase inhibitor or placebo. Unfortunately one patient appeared intolerant of the allocated treatment. For this patient the code was broken and the treatment was found to be the active drug; the patient was therefore removed from the trial. The 11 remaining patients successfully completed the four months of active treatment or placebo, followed by repeat neuroimaging.

The six controls underwent neuroimaging on one occasion only. Jack-knifing the QNB control studies (a procedure whereby each of the six subjects is removed in turn and compared with the remaining five subjects) within SPM99 showed that one of these control subjects had significant abnormalities. A subsequent rCBF SPECT cerebral perfusion study demonstrated significant parieto-temporal hypoperfusion suggestive of early Alzheimer’s disease. Consequently this control was withdrawn from the study and will be kept under medical review.

A significant association was noted between the initial ADAS-COG and the baseline 123I-QNB images such that greater impairment on psychometric testing was associated with reduced uptake of 123I-QNB (p<0.05). This finding was based on the 10 patients who completed brain imaging and had complete ADAS-COG assessment.

No significant group differences (placebo v controls) were noted on the changes in neuropsychometry, nor were any significant associations found between the sequential changes on psychometric assessment and brain imaging in this pilot study.

SPM99 analysis did not show any significant group differences between the baseline scans of the 12 patients and the five subjects in the control group.

A group comparison of the baseline 123I-QNB studies of the six patients on active treatment with the five patients randomised to placebo did not show any significant differences.

The abnormalities demonstrated on imaging for an individual patient showed very similar patterns between baseline and follow up studies for each patient, indicating that the 123I-QNB imaging technique is highly reproducible; however, there was considerable variability in the patterns of abnormalities between patients.

A paired t tests of the patients on active drug showed a reduction in tracer uptake in their four month follow up scan as compared with their baseline scan (311 voxels showed a significant decrease, p = 0.028; fig 1). Similarly, in the placebo group there was a reduction in the corresponding scans (745 voxels showed a decrease, p = 0.016; fig 2). For both these comparisons, up to 300 voxels would have been expected by chance.

A group comparison within SPM99 of the differences of the paired scans in the actively treated group (follow up study minus baseline) with the differences of the paired scans in the placebo group showed greater differences in the placebo group. These changes were centred on the left parieto-temporal region. By restricting the analysis to the posterior cortices (mild to moderate Alzheimer’s disease predominantly affects parieto-temporal grey matter), a significant cluster could be identified in the region of the precuneus in the left medial parietal lobe (p = 0.02).

DISCUSSION
In vivo imaging of the cerebral postsynaptic muscarinic receptor has been successfully undertaken using 123I-QNB, which appears to target the M1 muscarinic receptor preferentially; these receptors are predominantly located postsynaptically. Excellent correlations have been shown in healthy controls between the distributions of 123I-QNB uptake on in vivo imaging and muscarinic receptor densities in vitro.

Reduced M1 receptor uptake in patients with Alzheimer’s disease has been observed by others, predominantly in the parieto-temporal cortices. Although this is at variance with several histopathological findings of normal receptor density, the most likely explanation is that, although the...
Cholinergic treatment in Alzheimer’s disease

The M1 receptor and the G protein related signal non-functioning. A possible mechanism is uncoupling receptors may be structurally present at necropsy, they are compared with the first for the patient group receiving placebo. Note the areas of significant deterioration in the second scan as Figure 2. Cholinergic treatment in Alzheimer’s disease.15 16 An in vivo animal study using the positron emission tomography (PET) ligand 18F-FP-TZTP to target the M1 and M2 muscarinic receptors showed only a non-significant reduction in cortical ligand binding when acetylcholine concentrations were increased by the administration of the acetylcholinesterase inhibitor physostigmine.27 In our study, any displacement of the QNB ligand by raised endogenous levels of acetylcholine in the actively treated group would have underestimated the significant differences noted in receptor binding between the actively treated and placebo groups.

Conclusions

The results of this pilot study show that patients on cholinergic treatment have better preservation of M1 receptor binding to 123I-QNB than those receiving placebo. This suggests a possible neuroprotective role for cholinergic treatment in Alzheimer’s disease.

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Competing interests: Pfizer-Eisai have awarded a project grant to DW and CH, involved them in multicentre trials, and contributed to their expenses for conference presentations.

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Cerebral embolism in endocarditis: William Senhouse Kirkes (1823–64)

The distinction between thrombosis and haemorrhage was unclear until the mid-19th century,¹ despite the clinical and pathological descriptions of Abercrombie, Cheyne, Cooke, and, in France, Serres. Small softenings were first paper6 described together with the later paper of Samuel Wilks brought this published one of the earliest descriptions of cerebral T

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Reference


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