SHORT REPORT

Is there any evidence for a protective effect of antithrombotic medication on cognitive function in men at risk of cardiovascular disease? Some preliminary findings

M Richards, T W Meade, Sir Stanley Peart, P J Brennan, A H Mann

Abstract

To explore whether antithrombotic medication may protect against cognitive decline, tests of verbal memory, attention, abstract reasoning, verbal fluency, and mental flexibility were administered to 405 men at risk of cardiovascular disease. These subjects were a subgroup of those who had been participating in a randomised double blind factorial trial of low dose aspirin (75 mg daily) and low intensity oral anticoagulation with warfarin (international normalised ratio of 1.5) at 35 general practices across the United Kingdom for at least five years, were at least 55 years old at trial entry, and had been randomly allocated to one of four groups: active warfarin and active aspirin, active warfarin and placebo aspirin, placebo warfarin and active aspirin, and double placebo. Verbal fluency and mental flexibility were significantly better in subjects taking antithrombotic medication than in subjects taking placebo. Aspirin may have contributed more than warfarin to any beneficial effect. These results provide tentative evidence that antithrombotic medication may protect cognitive function in men at risk of cardiovascular disease.

antithrombotic medication on cognitive function in those without frank dementia. Population based studies have either reported no association² or a modest trend towards protection of cognitive function³ from medication (non-steroidal anti-inflammatory drugs and aspirin). However, these authors relied on recall of use of medication and, in both studies, employed a single, global measure of current cognitive status.

In the present study we measured cognitive function in patients already enrolled in a double blind placebo controlled factorial trial of low dose aspirin and low intensity oral anticoagulation with warfarin for their effect on the prevention of coronary and cerebral thrombosis. The purpose was to assess whether those who were on active treatment showed differences in cognitive performance compared with those who were receiving placebo. Because this cognitive study was conducted after the trial was already in progress, no baseline cognitive test data are available for comparison. Nevertheless, we were able to take advantage of the randomised design and detailed monitoring during a five year interval. The content of our neuropsychological test battery was based on studies that reported an association between cardiovascular risk factors and impairments in memory, attention, and abstract reasoning.⁴⁻⁶

Methods

MEDICAL RESEARCH COUNCIL THROMBOSIS PREVENTION TRIAL (TPT)

This trial is taking place within the Medical Research Council's general practice research framework and involves 101 collaborating general practices. A chart review of all men aged between 45 and 69 in these practices was carried out by the trial nurse. Those who were not excluded on grounds of contraindications to a TPT drug (aspirin and warfarin) or likely poor compliance, were invited to attend for screening to assess cardiovascular risk status. Screening procedures are described in detail elsewhere^{7,8} and identified those in the top quintile of risk for myocardial infarction or coronary death, using a score based on family and smoking history, blood pressure, body mass index, blood cholesterol, plasma fibrinogen, and plasma factor VII activity. Those at high risk were invited to enter the trial, in

Section of Old Age Psychiatry
M Richards

Section of Epidemiology and Primary Care, Institute of Psychiatry, London
A H Mann

MRC Epidemiology and Medical Care Unit, Wolfson Institute of Preventive Medicine, Medical College of St Bartholomew's Hospital, London, UK
T W Meade
S Peart
P J Brennan

Correspondence to:
Dr Marcus Richards,
Institute of Psychiatry, De Crespigny Park, Denmark Hill, London SE5 8AF, UK.

Received 22 April 1996
and in final revised form
25 November 1996

Accepted 29 November 1996

(*J Neurol Neurosurg Psychiatry* 1997;62:269-272)

Keywords: cognitive; aspirin; warfarin; cardiovascular; antithrombotic

A growing body of evidence suggests an association between cardiovascular risk factors and cognitive impairment during aging. These risk factors include hypertension, diabetes, hypercholesterolaemia, and cardiac disease such as atrial fibrillation and angina, all of which may promote neuropathological changes resulting from microthromboses in cerebral blood vessels. Antithrombotic medication may therefore protect cognitive function. In a randomised placebo controlled trial, Meyer *et al*¹ found that aspirin improved cognitive function in patients with multi-infarct dementia. Little is known, however, about the effect of

which they were randomised to active warfarin and active aspirin (WA), active warfarin and placebo aspirin (W), active aspirin and placebo warfarin (A), or double placebo treatment (P).

Participants are examined at entry and, once stabilised on treatment, at three monthly intervals by the trial nurse and at annual examinations by the general practitioner. Treatment with warfarin is initiated at a dose of 2.5 mg daily and increased by 0.5 or 1.0 mg steps until the intended international normalised ratio (INR) of about 1.5 has been reached. On average, this stabilisation process takes about four months. Dose changes are made equally often for those receiving placebo warfarin treatment with identical tablets. The dose of aspirin is 75 mg daily in a controlled release formulation, also with identical placebo tablets.

Subjects

Subjects eligible for this cognitive substudy were (1) aged at least 55 at entry and (2) had been participating in the trial for five years or more. Older trial entrants were thought to be most likely to show cognitive changes, and those participating in the trial for five years had received antithrombotic treatment for long enough for any effect of treatment on cognitive function to be evident. Of all 101 practices, 35 included men meeting the above criterion of treatment for at least five years. A difference of five seconds in part A of the trailmaking test (which has a large variance) is of potential functional significance for subjects in the age group of those in the present study.⁹ A sample of 350 subjects was estimated to be necessary to detect this effect with a power of 90% at a 5% level of significance. Of 1001 subjects originally meeting the above criteria, 493 had either already completed the TPT study or were no longer participating. Morbidity data, including causes of adverse events and death, will be available at the end of the TPT study. Of the remaining 508 eligible subjects, 60 were in practices where training for the neuropsychological tests could not be arranged and 24 subjects declined to be tested (nine in the P group, six in the A group, six in

the W group, and two in the WA group). Thus 424 subjects underwent cognitive testing. All subjects gave signed informed consent.

COGNITIVE MEASURES

The following tests were administered at the relevant follow up assessment but not at trial entry. (1) Logical memory subtest of the Wechsler memory scale¹⁰ (immediate and delayed); a measure of memory decline was obtained by subtracting the delayed from the immediate score and this measure was used in the present analyses. (2) Digit span subtest of the revised Wechsler adult intelligence scale (WAIS-R¹¹). (3) A cancellation task¹²; subjects were required to cross out, as quickly as possible, letter triads (TMX) from an array of similar triads. The lower the score, the better the performance. (4) Trailmaking test¹³ (parts A and B). A measure of mental flexibility was gained by subtracting part A from part B¹⁴ and this measure was used in the present analyses. The lower the value, the better the performance. (5) The similarities subtest of the WAIS-R was used to assess abstract reasoning. (6) Verbal fluency was assessed by (a) production of animal names¹⁵ and (b) production of words beginning with the letter F.¹⁶ One minute was allowed for each.

Five regional nurse coordinators underwent standardised training by the first author to administer the neuropsychological tests. They then instructed clinic nurses, who administered the tests blind to medication status. This training required observation and practice and emphasised strict adherence to formal written protocols for each test.

Results

BASELINE CHARACTERISTICS

Table 1 shows the baseline characteristics for subjects in the four treatment groups who underwent cognitive testing.

There were no significant differences in age, education, body mass index, systolic or diastolic blood pressure, number of cigarettes smoked per day, plasma cholesterol, plasma fibrinogen, ECG, ischaemia, or family history of heart disease. Nor was there any difference between the groups in duration in the trial at the time of cognitive testing. Subjects in the placebo group had a slightly higher factor VII activity level at baseline compared with subjects in the active treatment groups ($P = 0.04$). There were no significant differences in these baseline characteristics between subjects who underwent cognitive testing and those from the original pool of 1001 who would have been eligible for cognitive testing.

COGNITIVE FUNCTION AND ANTITHROMBOTIC MEDICATION

Subjects were first grouped into those receiving placebo only and those on any active medication (WA, A, or W). Table 2 shows the cognitive scores for these two groups.

Multivariate analysis of variance (MANOVA) was employed, using Hotelling's T^2 statistic to compare the multiple cognitive

Table 1 Baseline risk factor and background demographic data for the four treatment groups

Variable	Groups			
	WA	W	A	P
No	108	107	97	93
Age at testing (y)	66.7 (3.8)	66.9 (4.1)	66.0 (3.5)	66.1 (4.0)
Education (y)	10.0 (2.2)	10.0 (2.2)	10.7 (2.3)	10.1 (1.8)
Time in trial (y)	5.5 (1.0)	5.4 (0.9)	5.5 (0.9)	5.6 (1.2)
Body mass index (wt/ht ²)	38.1 (10.3)	38.4 (10.0)	37.8 (12.0)	40.3 (13.9)
Systolic blood pressure (mm Hg)	139.5 (17.6)	144.2 (18.2)	142.1 (19.5)	142.7 (18.5)
Diastolic blood pressure (mm Hg)	81.1 (10.0)	82.8 (9.4)	82.2 (9.7)	83.6 (11.6)
Cigarettes/day	4.6 (8.5)	5.6 (8.2)	6.7 (9.6)	4.5 (8.2)
Factor VII (% standard)	104.6 (26.6)	104.2 (27.3)	108.7 (28.5)	114.4 (28.2)
INR	1.0 (0.1)	1.0 (0.1)	1.0 (0.1)	1.0 (0.1)
Plasma cholesterol (mmol/l)	6.3 (1.3)	6.1 (1.2)	6.0 (1.8)	6.3 (1.6)
Plasma fibrinogen (g/l)	296.6 (82.3)	306.5 (93.6)	287.4 (104.1)	290.4 (83.3)
Family history of heart disease (%)	15.7	14.0	18.6	20.4
ECG ischaemia (%)	2.8	4.7	5.2	3.2

Values are means (SD). INR = international normalised ratio. See text for groups.

Table 2 Cognitive scores for subjects taking active treatment v placebo only

Variable	Any active treatment	Placebo
No	312	93
Logical memory decline (s)	2.2 (1.5)	2.4 (1.5)
Digit span	11.4 (2.2)	11.6 (2.5)
Cancellation (s)	61.9 (17.9)	62.3 (17.8)
Similarities	17.3 (4.5)	17.6 (4.2)
Animal naming	20.2 (5.4)	18.9 (4.9)*
Letter naming (F)	12.6 (4.8)	12.0 (4.8)
Trails B-A (s)	50.2 (31.0)	59.7 (37.4)*

Values are means (SD).

* $p < 0.05$.

measures across these two groups. Nineteen subjects had missing values for at least one cognitive measure, including four subjects who completed the trailmaking B test more rapidly than trailmaking A, indicating probable test administration error. As MANOVA rejects cases with missing data for any variable, the results for these subjects were omitted from the analysis, leaving 405 men in the study. MANOVA disclosed a significant medication effect for the cognitive scores (Hotelling's $T^2 = 0.04$, $P = 0.013$). Corresponding univariate F tests disclosed poorer performance for subjects taking placebo only for animal naming ($F = 4.51$, $P = 0.034$) and trailmaking part B-A ($F = 6.06$, $P = 0.014$).

EFFECT OF SPECIFIC REGIMES

MANOVA was then used to compare cognitive function in the four separate treatment groups. Univariate F tests showed a significant group effect for WAIS-R Similarities ($F = 2.64$, $P = 0.049$), with poorest performance in the W group. However, as Hotelling's T^2 test was not significant, the possibility that this was a type I error cannot be ruled out.

These analyses suggested the possibility of better cognitive performance by patients taking aspirin than in patients taking warfarin. Table 3 therefore shows the results for all those taking aspirin (WA and A) compared with those not taking aspirin (W and P) and for all those taking warfarin (WA and W) compared with those not taking warfarin (A and P)—that is, a factorial design for separate treatment effects.

In the comparison according to warfarin allocation, several test scores were, if anything, poorer in those on active warfarin than with those who were not, significantly so for similarities ($F = 5.49$, $P = 0.02$), although, once again, Hotelling's T^2 test was not significant.

EFFECT OF RISK FACTOR VARIABLES

Correlations were calculated between time to complete the trailmaking B-A test and the cardiovascular risk factor values. Values at screening (see methods), as close as possible to the time of cognitive testing and the mean across the repeated assessments were used. All correlation coefficients fell within a range of -0.1 to 0.1 . Thus there were virtually no associations between the risk factors and the trailmaking B-A score. A similar pattern was observed when the P group and the combined treatment groups were analysed separately. There was no significant difference in the trailmaking B-A score between subjects with and without a family history of cardiovascular disease.

Discussion

As part of a double blind factorial trial of the effect of antithrombotic medication on the prevention of coronary heart disease we found that 405 of the older subjects who had been taking low dose or low intensity antithrombotic medication with warfarin, or aspirin, or both for about five years showed better cognitive performance than subjects given only placebo. We found this trend in most cognitive scores. Any effect was probably mainly due to aspirin. More specifically, scores for animal naming (a test of verbal fluency) and part B-A of the trailmaking test were significantly better in subjects taking active medication. Both of these tests are effortful, require mental control and flexibility, and are thought to be under frontal control. The trailmaking tests require motor speed and visual scanning although these elements were controlled in part by subtracting part A from part B and were not, in any case, sensitive to medication in the cancellation test.

Caution is necessary in interpreting the results of this or any other study that consists of a subgroup analysis within a trial that was designed for another purpose. Firstly, we did not measure cognitive function at baseline. The possibility that treatment might influence cognitive function was only raised after the trial had started. We have to assume that the groups were similar in cognitive function at baseline, which seems likely. They were well matched in other respects and there was no discernable difference between those cognitively tested and those potentially available but not tested. Secondly, cognitive tests were administered by nurses with no previous experience in neuropsychological testing. Nevertheless, they received structured training, were guided by written test protocols, and were in any case blind to the medication status of each patient. Thirdly, the effect of antithrombotic medication on cognitive function, although significant, was generally slight although we would argue, in line with Gorkin *et al.*,⁹ that the effect for Part B-A of the trailmaking test (a difference of 9.5 seconds between placebo and active treatment groups) is of functional significance.

Whereas it should be born in mind that

Table 3 Cognitive scores for subjects taking or not taking aspirin and for subjects taking or not taking warfarin

	Taking aspirin		Taking warfarin	
	Yes (WA + A)	No (W + P)	Yes (WA + W)	No (A + P)
LM (decay)	2.2 (1.5)	2.2 (1.5)	2.1 (1.5)	2.4 (1.4)
Digit span	11.5 (2.3)	11.5 (2.4)	11.4 (2.2)	11.6 (2.4)
Cancellation (s)	61.3 (17.7)	62.7 (18.0)	62.0 (19.4)	61.9 (16.0)
Similarities	17.7 (4.2)	17.0 (4.7)	16.9 (4.7)	17.9 (4.0)
Animal naming	20.4 (5.4)	19.4 (5.1)	20.0 (5.3)	19.8 (5.3)
Letter naming (F)	12.6 (4.8)	12.3 (4.7)	12.4 (4.7)	12.5 (4.9)
Trails B-A (s)	49.1 (29.9)	55.8 (35.3)	51.8 (31.8)	53.1 (34.0)

Values are means (SD).

LM = Logical memory. See text for groups.

variance for the cardiovascular risk factors was restricted in this sample, the striking absence of correlations between any of the risk factors and time to complete part B-A of the trailmaking test was unexpected and brings into question the starting hypothesis of a vascular explanation for the wide range of scores found. Aspirin in the thrombosis prevention trial is low dose, 75 mg daily, and in a controlled release formulation designed for a prehepatic effect by reducing thromboxane in the platelets themselves while sparing prostacyclin production in the systemic vasculature after the liver metabolism of aspirin to salicylate, which is only weakly active.¹⁷ If aspirin does preserve cognitive function, it may perhaps do so through some pathway other than its effect on platelets and thrombogenesis, in which case a higher dose than the 75 mg used in the thrombosis prevention trial and in a formulation subject to less extensive first pass metabolism in the liver might result in a larger benefit. Further investigations of the association between antithrombotic medication and cognitive function are warranted.

The salary of MR is currently provided by the Alzheimer's Disease Society. We wish to acknowledge the help of the MRC general practice research framework doctors and nurses. Specific thanks are due to Mrs Pam Allen, Mrs Leslie Hand, Mrs Sue Fox, Mrs Margaret Goldsborough, and Mrs Eileen Marshall for coordination of nurse training for this study.

- 1 Meyer JS, Rogers RL, McClintic K, Mortel KF, Lofti J. Randomized clinical trial of daily aspirin therapy in multi-infarct dementia: a pilot study. *J Am Geriatr Soc* 1989;37:549-55.
- 2 May FE, Moore MT, Stewart RB, Hale WE. Lack of association of non-steroidal anti-inflammatory drug use and

- cognitive decline in the elderly. *Gerontology* 1992;38:275-9.
- 3 Sturmer T, Glynn RJ, Field TS, Taylor JO, Hennekens CH. Aspirin use and cognitive function in the elderly. *Am J Epidemiol* 1996;143:683-91.
- 4 Waldstein SR, Manuck SB, Ryan CM, Muldoon MF. Neuropsychological correlates of hypertension: review and methodologic considerations. *Psychol Bull* 1991;110:451-68.
- 5 Bornstein RA, Kelly MP. Risk factors for stroke and neuropsychological performance. In: Bornstein RA, Brown GG, eds. *Neurobehavioral aspects of cerebrovascular disease*. New York: Oxford University Press, 1991:182-201.
- 6 Desmond DW, Tatemichi TK, Paik M, Stern Y. Risk factors for cerebrovascular disease as correlates of cognitive function in a stroke-free cohort. *Arch Neurol* 1993;50:162-6.
- 7 Meade TW, Wilkes HC, Brennan PJ, Kelleher C, Browne W. Randomised controlled trial of low dose warfarin in the primary prevention of IHD in men at high risk: design and pilot study. *Eur Heart J* 1988;9:836-43.
- 8 Meade TW, Roderick PJ, Brennan PJ, Wilkes HC, Kelleher CC. Extracranial bleeding and other symptoms due to low dose aspirin and low intensity oral anticoagulation. *Thromb Haemost* 1992;68:1-6.
- 9 Gorkin L, Norvell NK, Rosen RC, Charles E, Shumaker SA, McIntyre KM, et al. Assessment of quality of life as observed from the baseline data of the studies of left ventricular dysfunction (SOLVD) trial quality-of-life sub-study. *Am J Cardiol* 1993;71:1069-73.
- 10 Wechsler D. A standardized memory scale for clinical use. *J Psychol* 1945;19:87-95.
- 11 Wechsler D. *Wechsler adult intelligence scale-revised*. New York: Psychological Corporation, 1981.
- 12 Reitan RM, Wolfson D. *The Halstead-Reitan neuropsychological test battery: theory and clinical interpretation*. Tuscon, AZ: Neuropsychology Press, 1985.
- 13 Taylor AE, Saint-Cyr JA, Lang AE. Frontal lobe dysfunction in Parkinson's disease. *Brain* 1986;109:845-83.
- 14 Sano M, Rosen W, Mayeux R. *Attention deficits in Alzheimer's disease*. Washington, DC: American Psychiatric Association, 1984.
- 15 Goodglass H, Kaplan D. *The assessment of aphasia and related disorders*. 2nd ed. Philadelphia, PA: Lea and Febinger, 1983.
- 16 Benton AL, Hamsher K. *Multilingual aphasia examination*. Iowa City: University of Iowa Hospitals, 1978.
- 17 Clarke RJ, Mayo G, Price P, Fitzgerald GA. Suppression of thromboxane A2 but not of prostacyclin by controlled-release aspirin. *N Engl J Med* 1991;325:1137-41.