Renin angiotensin system inhibition is associated with reduced free radical concentrations in arteries of patients with coronary heart disease

Angiotensin II, which is also thought to play a key role in atherosclerosis, has recently been shown to have pro-oxidant effects, by increasing superoxide (O$_2^−$) production in human arteries. Oxidative stress, a state of excessive free radical activity which is associated with reduced bioavailable nitric oxide (NO), may be evident in patients with coronary heart disease (CHD). Lately ACE inhibitors have been shown to reduce cardiovascular morbidity and mortality in patients at high risk of CHD. The aim of the present study was to determine, which, if any, risk factors and drug treatments were associated with altered free radical concentrations in the arteries of CHD patients undergoing coronary artery bypass grafting (CABG).

Seventy-nine consecutive patients who were undergoing CABG were prospectively included in this study. Patient characteristics were determined by review of case records. A history of current cigarette smoking, hypertension (defined as either current antihypertensive treatment or a blood pressure > 140/90 mm Hg), diabetes mellitus, and hypercholesterolaemia (plasma cholesterol > 5.5 mmol/l) were considered as risk factors for CHD.

Distal segments of left internal mammary artery which were obtained at the time of CABG were taken to the laboratory in Krebs-Hepes buffer (pH 7.4 ± 0.2), carefully dissected free of loose connective tissue, divided into 4-5 mm segments and weighed. Vascular O$_2^−$ was measured by lucigenin enhanced chemiluminescence in a liquid scintillation counter (Hewlett Packard Model Tricarb 2100TR). Absolute counts were quantified with a xanthine/xanthine oxidase calibration curve for O$_2^−$ generation and reported as picomol per milligram per minute of tissue. Statistical analyses of vascular O$_2^−$ concentrations after log transformation were undertaken using the non-parametric Mann-Whitney Test and a stepwise multiple regression analysis was also performed. A probability value of $p < 0.05$ was considered significant. This study was approved by the hospital's ethics committee.

Data on age, sex, risk factors, and drug treatment are given in table 1. The profiles of risk factors and different classes of drug treatments were similar between patients who were taking an angiotensin converting enzyme (ACE) inhibitor or an angiotensin type 1 receptor antagonist (ARA), compared to those who were not taking these treatments. An ACE inhibitor or ARA was prescribed in 16 and three patients, respectively. The median rate of production of O$_2^−$ in internal mammary arteries was 1137 (interquartile range (IQR) 1290) pmol/mg/min. Superoxide concentrations were lower in those patients taking either an ACE inhibitor or an ARA (857 (IQR 670) pmol/mg/min; n = 19) compared to those who were not (1600 (IQR 1511) pmol/mg/min; n = 60; $p = 0.002$; 95% confidence interval for median difference 487 to 1228 pmol/mg/min (fig 1)). No other associations between age, sex, risk factors or drug treatments and superoxide concentrations were identified.

We have shown, for the first time in human arteries, that O$_2^−$ concentrations were lower in patients treated with either an ACE inhibitor or an ARA, compared to those who were not. The clinical characteristics of both of these groups were similar such that the observed differences in vascular O$_2^−$ concentrations were unlikely to be explained by any other patient characteristic or treatment. It is of interest that despite the fact that a large proportion of these patients were taking drugs with putative antioxidant properties, such as aspirin, β blockers and HMG-CoA reductase inhibitors, vascular free radical concentrations were detected at physiologically important concentrations. The absence of any important antioxidant effect of these other treatments suggests that the sample size may not be sufficiently large to detect what may be a lesser antioxidant effect of these drugs. The variation in basal vascular O$_2^−$ concentrations observed in this and other investigations in humans, and the lack of correlation of O$_2^−$ production with some atherosclerotic risk factors, may be caused by the heterogeneous clinical characteristics of patients with CAD.

Our observations raise two questions. The first is how does such treatment exert this effect? Though some ACE inhibitors may have direct free radical scavenging properties, this effect has been difficult to show at therapeutic concentrations in humans. A more likely explanation is that the antioxidant effect of this treatment is caused by inhibition of the effects of angiotensin II. The second question is what, if any, might be the therapeutic significance of this effect of RAS inhibition? A reduction in vascular free radical production associated with RAS inhibition, as is the case in the current study, may lead to enhanced bioavailable nitric oxide in vivo.

Taken together, these observations suggest that RAS inhibition leads to a reduction in oxidative stress in patients with CHD. Given the damaging effects of increased free radical activity in the vasculature, the antioxidant effects of these treatment may be one further mechanism which may contribute to their beneficial effects in patients with CHD.

COLIN BERRY
NIALL ANDERSON
ALAN J B KIRK*
ANNA F DOMINICZAK
JOHN J V MCMURRAY
*Department of Cardiothoracic Surgery,
North Glasgow Hospitals University Trust,
Western Infirmary, Glasgow, UK

Correspondence to: Dr Colin Berry, Department of Medicine and Therapeutics, University of Glasgow, 44 Church Street, Glasgow G11 6NT, UK;
colin.berry@clinmed.gla.ac.uk

Sources of support: CB is a Medical Research Council Clinical Training Fellow. This work is also supported by a British Heart Foundation Programme Grant (RG/09009) to AFD.

www.heartjnl.com
Survival is shown as actual percentages at 3 years and as Kaplan-Meier (K-M) estimates at 6.1 years. The p values test the hypothesis that 6.1 year survival in a group equals survival in the age matched reference group with a normal ECG and N-ANP.

N-ANP concentrations, abnormal ECG, and dysfunction in heart patients from general practice remains to be defined. N-terminal atrial natriuretic peptide (BNP) are associated with heart patients from general practice.

Table 1  Survival of patients according to having an ECG with QRS or ST changes and N-terminal atrial natriuretic peptide (N-ANP) > 0.8 nmol/l or not

<table>
<thead>
<tr>
<th>Age group</th>
<th>Total (n)</th>
<th>Deaths (n)</th>
<th>Survival actual (%)</th>
<th>3 year follow up</th>
<th>Survival K-M (%)</th>
<th>6.1 year follow up</th>
<th>Median survival (years)</th>
<th>Log rank test p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients &lt; 70 years of age (n=83)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal ECG, normal N-ANP</td>
<td>26</td>
<td>1</td>
<td>96</td>
<td>96</td>
<td>&gt; 6.1</td>
<td>&gt; 6.1</td>
<td>Reference</td>
<td>0.052</td>
</tr>
<tr>
<td>Abnormal ECG, normal N-ANP</td>
<td>22</td>
<td>2</td>
<td>91</td>
<td>84</td>
<td>&gt; 6.1</td>
<td>&gt; 6.1</td>
<td>Reference</td>
<td>0.052</td>
</tr>
<tr>
<td>Normal ECG, raised N-ANP</td>
<td>1</td>
<td>0</td>
<td>100</td>
<td>100</td>
<td>&gt; 6.1</td>
<td>&gt; 6.1</td>
<td>Reference</td>
<td>0.052</td>
</tr>
<tr>
<td>Abnormal ECG, raised N-ANP</td>
<td>4</td>
<td>3</td>
<td>25</td>
<td>25</td>
<td>1.6</td>
<td>&lt; 0.0001</td>
<td>Reference</td>
<td>0.052</td>
</tr>
</tbody>
</table>

The p values test the hypothesis that 6.1 year survival in a group equals survival in the age matched reference group with a normal ECG and N-ANP. Survival is shown as actual percentages at 3 years and as Kaplan-Meier (K-M) estimates at 6.1 years.
study. Nor have we evaluated those variables in acute patients, which is a different situation." Just half the target group accepted our invitation for a full cardiac assessment. Those who withdrew were old, they were probably more disabled, and their prognosis may have been worse than for subjects who were examined. In view of the understandable withdrawal pattern, we believe our results are applicable to a scenario in general practice where a physician wants to make a risk assessment for a presumed heart patient without overt congestive heart failure. A normal ECG will convince the physician and patient of a favourable prognosis and a low likelihood of left ventricular systolic dysfunction. Our study suggests that measurement of natriuretic peptide is particularly useful in subjects with an abnormal ECG. If both N-ANP and the ECG are abnormal, further cardiac assessment seems sensible to determine whether treatable conditions exist, although we have no data on whether cardiac assessment will indeed change the outcome.

OLAV WENDELBOE NIELSEN
JØRGEN HILDEN*
JØRGEN FISCHER HANSEN
Cardiovascular Department,
Copenhagen University Hospital Bispebjerg,
DK-2400 Bispebjerg, Denmark
*Department of Biostatistics,
University of Copenhagen,
Copenhagen, Denmark

Correspondence to: Dr Nielsen, Måkhærg Alle
66, 2970 Hørsholm, Denmark;
OWN@dadlnet.dk

Oliv Wendelboe Nielsen formed the primary study hypothesis and core ideas, designed the protocol, made the data analysis and wrote the paper. Jørgen Fischer Hansen discussed core ideas and participated in writing of the paper. Jørgen Hilden participated in statistical analysis and the writing of the paper. OWN will act as guarantor of the paper. The study was supported by the Danish Heart Foundation.


Doxetaxel induced cardiotoxicity

Doxetaxel is a new taxoid antineoplastic agent.1 Its mechanism of action is primarily related to its ability to enhance microtubule assembly and to stabilize microtubules by preventing their depolymerisation, thus disrupting normal cell division.2 Since doxetaxel has significant cytotoxic activity against human breast cancer in vitro and in vivo, it is widely used in patients with breast cancer, especially those with metastatic breast cancer. It has been shown that increased microtubule density, for which microtubule stabilisation is one potential mechanism, causes contractile dysfunction in cardiac hypertrophy.3 Since doxetaxel exerts its actions by stabilising microtubules, it is reasonable to consider that doxetaxel may induce contractile dysfunction as a cardiotoxic agent. Thus, we examined the effect of doxetaxel on cardiac function and the serum concentration of brain natriuretic peptide (BNP) and other neurohormones one day before, day one after, and three weeks after doxetaxel administration.

Ten consecutive patients with breast cancer having skin metastasis who received doxetaxel were investigated. All patients were women and their mean (SEM) age was 51.8 (2.5) years. Doxetaxel dissolved into 500 ml of 0.9% saline solution was administered with an intravenous drip injection for three hours. The mean dosage of doxetaxel was 52.36 (6.35) mg/m2. Although the serum concentration of brain natriuretic peptide (BNP) was within normal range before treatment, it was significantly raised on the day after doxetaxel administration (table 1). In contrast, concentrations of other circulating cardiac neurohormones (atrial natriuretic peptide, renin, aldosterone, noradrenaline (norepinephrine), and adrenaline (epinephrine)) were within the normal range at both determinative points (table 1). Additionally, blood pressure was not affected by this treatment (one day before chemotherapy 126 (7)/74 (6) mm Hg; one day after chemotherapy 123 (8)/76 (8) mm Hg; three weeks after chemotherapy 127 (5)/69 (9) mm Hg; NS). To assess left ventricular systolic and diastolic functions, we measured fractional shortening, ejection fraction, mitral inflow E:A ratio, and deceleration time by echocardiography. While fractional shortening and ejection fraction were not affected by doxetaxel, the E:A ratio was significantly decreased and deceleration time was significantly increased (table 1). Additionally, the maximum diameter of inferior vena cava was not affected by this treatment. These results suggest that doxetaxel induced left ventricular diastolic dysfunction and an increase in serum BNP concentration without increasing preload or afterload.

BNP is a useful biochemical marker of left ventricular dysfunction, having diagnostic, therapeutic, and prognostic implications.4 In addition, diastolic dysfunction is well known as a cause of congestive heart failure.5 In this study, the increase in BNP concentration, the decrease in E:A ratio, and the increase in deceleration time were transient and returned to baseline within three weeks in all patients (table 1). However, since doxetaxel induced the transient abnormalities of the serum BNP concentration, E:A ratio, and deceleration time, even in patients with normal cardiac function, it is possible that this agent may induce heart failure in patients with left ventricular diastolic dysfunction. Consequently, it is advisable to check the cardiac function of patients to be treated with doxetaxel by echocardiography. In addition, serum BNP can be a useful marker by which to monitor doxetaxel induced cardiotoxicity.


Implementation of the NICE guidelines for the primary prevention of mortality from ventricular tachyarrhythmias: implications for UK electrophysiology centres; activity modelling from the UK-HEART study

It is now well established that the implantable cardioverter-defibrillator (ICD) is the most effective treatment for the primary prevention of life threatening ventricular arrhythmia.6 7 Despite this, its widespread use in the UK for the indication has been minimal, at least in part because of perceived resource implications. In September 2000, the National Institute for Clinical Excellence (NICE) recommended electrophysiological testing (EPS) for all patients with a history of myocardial infarction who have an ejection fraction of less than 35% and three or more beats of non-sustained ventricular tachycardia (NSVT) on a 24 hour Holter monitor. NICE further recommended that all patients in whom a significant ventricular arrhythmia is induced at EPS should have an ICD

Table 1 Serial changes in biochemical and echocardiographic parameters

<table>
<thead>
<tr>
<th>Parameter</th>
<th>1 day before chemotherapy</th>
<th>1 day after chemotherapy</th>
<th>3 weeks after chemotherapy</th>
<th>Reference values</th>
</tr>
</thead>
<tbody>
<tr>
<td>BNP (pg/ml)</td>
<td>13.7 (1.9)</td>
<td>53.8 (3.5)*</td>
<td>12.6 (2.4)</td>
<td>&lt; 20</td>
</tr>
<tr>
<td>ANP (pg/ml)</td>
<td>24.9 (4.9)</td>
<td>30.3 (4.6)</td>
<td>27.4 (3.9)</td>
<td>&lt; 40</td>
</tr>
<tr>
<td>Noradrenaline (pg/ml)</td>
<td>277.1 (35.3)</td>
<td>277.1 (33.8)</td>
<td>257.9 (51.9)</td>
<td>100–450</td>
</tr>
<tr>
<td>Adrenaline (pg/ml)</td>
<td>33.0 (6.0)</td>
<td>32.4 (5.7)</td>
<td>39.0 (12.8)</td>
<td>&lt; 100</td>
</tr>
<tr>
<td>Renin (resting state) (ng/ml/h)</td>
<td>1.37 (0.23)</td>
<td>1.30 (0.23)</td>
<td>1.09 (0.68)</td>
<td>0.2–2.7</td>
</tr>
<tr>
<td>Aldosterone (pm/ml)</td>
<td>70.6 (8.2)</td>
<td>69.2 (7.0)</td>
<td>74.6 (16.0)</td>
<td>45–105.5</td>
</tr>
<tr>
<td>Fractional shortening (%)</td>
<td>36.4 (1.0)</td>
<td>36.0 (1.2)</td>
<td>37.5 (0.9)</td>
<td>&gt; 28</td>
</tr>
<tr>
<td>Ejection fraction (%)</td>
<td>72.9 (5.2)</td>
<td>73.2 (6.3)</td>
<td>72.4 (7.4)</td>
<td>&gt; 56</td>
</tr>
<tr>
<td>Mitral inflow E:A ratio</td>
<td>1.45 (0.16)</td>
<td>0.71 (0.24)*</td>
<td>1.29 (0.12)</td>
<td>&gt; 1.0</td>
</tr>
<tr>
<td>Deceleration time (ms)</td>
<td>192.1 (9.9)</td>
<td>271.3 (16.3)*</td>
<td>209.2 (13.6)</td>
<td>199 (32)</td>
</tr>
<tr>
<td>Maximum diameter of IVC (mm)</td>
<td>15.1 (1.6)</td>
<td>12.9 (1.0)</td>
<td>12.0 (2.0)</td>
<td>&lt; 23</td>
</tr>
</tbody>
</table>

All data are expressed as mean (SEM).

Two way analysis of variance (ANOVA) and Fisher’s exact test for post hoc analysis were carried out for multiple comparisons among groups. *p < 0.05 versus a day before chemotherapy.

Reference


www.heartjnl.com

Downloaded from http://heart.bmj.com/ on June 22, 2017 - Published by group.bmj.com
implanted. The potential effect of the NICE guidelines on workload in UK electrophysiology centres has not been evaluated.

UK-HEART examined the utility of heart rate variability as a predictor of mortality in heart failure patients. Patients with stable heart failure were enrolled in four UK cardiac centres over a 17 month period (1 December 1993 to 30 April 1995). Patients with diabetes and atrial fibrillation were excluded; otherwise allcomers to cardiology clinics were eligible. All patients underwent echocardiographic estimation of ejection fraction and screening for arrhythmia, including NSVT, using 24 hour ambulatory electrocardiography. The baseline demographic and clinical characteristics of these patients were similar to those in MADIT® and MUSTT, the two primary prevention trials on which the NICE guidelines are based.

A total of 535 patients were entered into UK-HEART, of whom 551 had interpretable 24 hour ECGs. Of these, 57 fulfilled the NICE criteria for EPS. If diabetics and patients with atrial fibrillation had been included this figure would increase to 88 as approximately 35% of heart failure populations suffer from either of these conditions. Recent data have demonstrated that a second 24 hour tape will identify a further 38% of patients with NSVT, increasing numbers to 142. In MUSTT, sustained ventricular arrhythmia was induced at EPS in 35% of patients. Other smaller studies in similar populations have found equivalent positivity rates. If a cohort reacted in the same way there would be 50 positive studies and hence 50 ICD implants.

During the period of recruitment 68 electrophysiological studies for ventricular arrhythmia were performed on patients within the four centres. During an equivalent, contemporary period (1998-2000), 72 studies were performed. Application of NICE guidelines would lead to a further 142 studies, equivalent to three extra per month in each of the three centres performing this procedure. During 1993-95 there were 11 ICDs implanted. Between 1998-2000 there were 79 implanted. The NICE guidelines would mean a further increase of 50, representing one extra ICD per implanting centre per month.

The UK-HEART study was primarily designed to investigate the prognostic utility of measures of heart rate variability in a heart failure population and therefore has some limitations in its application to NSVT. Diabetics and patients with atrial fibrillation were excluded from UK-HEART because of the heart rate variability analysis. Diabetics make up 20% and patients with atrial fibrillation 15% of major heart failure studies. We have assumed that if these patients had been included, our population eligible for further investigation would increase by 35%. Clearly there will be some crossover between the groups and they may have different rates of NSVT. The patients in the study were allcomers to cardiology clinics. Numbers would increase if all primary and secondary care physicians had enrolled patients.

The numbers of investigations and ICDs estimated apply to the first year of guideline implementation. The numbers requiring investigation and treatment the following year, including new diagnoses and those whose disease had progressed, would differ; theoretically fewer than in the first year. The numbers would also fall as some patients eligible for investigation or treatment would be unsuitable for non-cardiological reasons.

Overall, these data provide evidence that implementation of the NICE guidelines for the use of ICDs in primary prevention is unlikely to lead to an unmanageable increase in EPS or ICD implantation in UK electrophysiology centres.

Correspondence to: NP Gall, Department of Cardiology, King’s College Hospital, Denmark Hill, London SE5 9RS, UK; nick.gall@kcl.ac.uk

**WEB TOP 10**

www.heartjnl.com

These articles scored the most hits on Heart’s web site during May 2001

1 Correlation between high frequency intravascular ultrasound and histomorphology in human coronary arteries
   P Prati, E Arbustini, A Labellarte, B Dal Bello, L Sommariva, MT Maltus, A Pagano, A Boccanelli
   May 2001;85:567–70. (Basic research)

2 Joint British recommendations on prevention of coronary heart disease in clinical practice
   December 1998;80(suppl 2):S1–29

3 Guideline for the management of patients with acute coronary syndromes without persistent ECG ST segment elevation
   February 2001;85:133–42

4 National Institute for Clinical Excellence guidance: too NICE to glycoprotein IIb/IIIa inhibitors?
   CJ Knight
   May 2001;85:481–3. (Editorial)

5 Acute myocardial infarction: primary angioplasty
   F Zigrara
   June 2001;7:705–9. (Education in Heart)

6 Cost effectiveness of ranipril treatment for cardiovascular risk reduction
   IS Malik, VK Bhatia, J S Kooner
   May 2001;85:539–43. (Cardiovascular medicine)

7 Anatomic basis of cross-sectional echocardiography
   RH Anderson, SY Ho, SJ Brecher
   June 2001;85:716–20. (Education in Heart)

8 Aspirin for primary prevention of coronary heart disease: safety and absolute benefit related to coronary risk derived from meta-analysis of randomised trials
   PS Sarnamanganath, P Ghahramani, PR Jackson, EF Wallis, LE Ramsay
   March 2001;85:265–71. (Cardiovascular medicine)

9 Absolute, attributable, and relative risk in the management of coronary heart disease
   JEC Sedgwick
   May 2001;85:491–2. (Editorial)

10 The fate of acute myocarditis between spontaneous improvement and evolution to dilated cardiomyopathy: a review
   A D’Ambrosio, G Patti, A Manzoli, G Sinagra, A Di Lenarda, F Silvestri, G Di Sciascio
   May 2001;85:499–504. (Review)

Visit the Heart website for hyperlinks to these articles, by clicking on “Top 10 papers”

www.heartjnl.com
Renin angiotensin system inhibition is associated with reduced free radical concentrations in arteries of patients with coronary heart disease

COLIN BERRY, NIALL ANDERSON, ALAN J B KIRK, ANNA F DOMINICZAK and JOHN J V MCMURRAY

Heart 2001 86: 217-220
doi: 10.1136/heart.86.2.217

Updated information and services can be found at:
http://heart.bmj.com/content/86/2/217

These include:

References
This article cites 5 articles, 2 of which you can access for free at:
http://heart.bmj.com/content/86/2/217#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections

- Hypertension (3006)
- Epidemiology (3752)
- Interventional cardiology (2933)
- Drugs: cardiovascular system (8842)
- Diabetes (842)
- Metabolic disorders (1030)
- Tobacco use (635)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/