The use of angiotensin-converting enzyme inhibitors in the treatment of heart failure in hospital practice

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Summary
Several well-controlled trials in patients with heart failure have shown that the use of angiotensin-converting enzyme (ACE) inhibitors, in combination with a diuretic, causes a reduction in mortality and morbidity, which seems to be mainly due to a reduction in fatal and nonfatal cardiovascular events. Our aim was to determine whether 249 consecutive patients discharged from hospital with a primary diagnosis of heart failure were routinely being treated with an ACE inhibitor at an appropriate dose. At the time of admission to hospital, 91 (36.5%) were receiving a combination of a diuretic and an ACE inhibitor, 129 (51.8%) were receiving a diuretic alone, and 29 (11.6%) had not previously received either a diuretic or an ACE inhibitor. At the time of discharge from hospital all patients were on a diuretic and 144 (57.8%) were also receiving an ACE inhibitor. Although 41 patients (16.5%) had a relative or absolute contraindication for the use of an ACE inhibitor, 64 patients (25.7%) with no contraindication were not receiving an ACE inhibitor. Many of the patients who were prescribed an ACE inhibitor were given it at an inappropriate dose; only 24 patients (16.7%) were on the dose that was used in the clinical trials showing a reduction in mortality. These results show that in one in four patients admitted to hospital with heart failure who should be receiving an ACE inhibitor by the time of discharge, are not. The average age of these patients was 76 years. Whilst it has been shown that the benefit of ACE inhibitors does not appear to be age-related, most published studies have not included many patients over the age of 80. Specific studies looking at the effect of ACE inhibitors in elderly patients would be helpful, as well as studies to determine the optimum treatment regimen for this age group.

Keywords: heart failure, ACE inhibitors

Congestive heart failure is a syndrome characterised by changes in left ventricular systolic and/or diastolic function which can ultimately result in reduced exercise tolerance, impaired quality of life, and a five-year mortality in excess of 50%. Heart failure imposes a heavy financial burden on any healthcare system due to the large number of hospital admissions. Recently, a series of randomised, double-blind controlled trials have consistently shown that, in patients already on a diuretic, the addition of an angiotensin-converting enzyme (ACE) inhibitor significantly improves symptoms and signs of chronic heart failure and increases survival. In addition, ACE inhibitors have been shown to slow or reverse left ventricular dilatation in patients with asymptomatic left ventricular dysfunction and attenuate left ventricular dilatation and progression to symptomatic heart failure in patients following myocardial infarction.

In a small pilot study in our hospital we found that some patients discharged from our institution with the diagnosis of heart failure were not receiving an ACE inhibitor. We therefore reviewed the notes of in-patients admitted to our hospital with heart failure to see how many were receiving ACE inhibitors and whether those who were not had any specific contraindication.

Patients and methods
We performed a retrospective analysis of all case notes of patients admitted to St George's Hospital during the period January to October 1994 with the primary diagnosis of heart failure. The diagnosis of heart failure was made by the admitting team based on clinical symptoms and signs and evidence of pulmonary oedema and/or cardiomegaly on a chest X-ray. In some patients the diagnosis of heart failure was supported by echocardiography. A list of all admissions was provided by the clinical coding department; 257 notes were requested, of which 249 were available. The results are expressed as means plus or minus standard deviations.

Results
A total of 249 subjects were included in the study (128 males; mean age 76 ±11 years). Two hundred and seven were white, 23 black and 19 Asian. One hundred and sixty were drawn from general medical firms, and the rest from the departments of geriatrics (66) and cardiology (23). The aetiology of heart failure, as stated in the notes, is shown in box 1. The
Aetiology of heart failure (n = 249)

<table>
<thead>
<tr>
<th>Aetiology</th>
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<tbody>
<tr>
<td>ischaemic heart disease</td>
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<tr>
<td>acute myocardial infarction</td>
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<tr>
<td>aortic and/or mitral valve disease</td>
<td>18</td>
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<tr>
<td>dilated cardiomyopathy</td>
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<td>fast atrial fibrillation</td>
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<td>hypertensive heart failure</td>
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<tr>
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Box 1

Diagnosis was confirmed in 146 patients (58.6%) using transthoracic echocardiography. Furthermore, 60 patients were on treatment for hypertension, 42 were diabetics (six on insulin), 21 had a cerebrovascular event, 13 had symptomatic peripheral vascular disease and 10 had renal impairment (plasma creatinine > 500 μmol/l).

At the time of admission to hospital of the 249 patients with heart failure, 91 patients (36.5%) were on a combination of a diuretic and an ACE inhibitor, 129 patients (51.8%) were receiving a diuretic alone and 29 (11.6%) had not previously received either a diuretic or an ACE inhibitor. At the time of discharge from hospital all patients were on a diuretic, 144 (57.8%) were also receiving an ACE inhibitor (67>75 years and 34>80 years of age). Forty-one patients (16.5%) had a relative or absolute contraindication for the use of an ACE inhibitor (box 2), but 64 (25.7%) patients with no contraindications to ACE inhibitors were not receiving them (51>75 years and 37>80 years of age).

In those patients receiving an ACE inhibitor, 92 patients were receiving treatment with captopril (mean daily dose 42 mg ± 25 mg); six were prescribed it once a day, 53 twice a day and 33 three times a day. Forty-eight patients were treated with enalapril (mean daily dose 17 mg ± 15 mg); 26 were prescribed it once a day, 21 twice a day and one three times a day. Four patients were on lisinopril (mean daily dose 14 mg ± 8 mg); all were on it once a day. Only 24 patients (16.7%) were on the dose that was used in the clinical trials where ACE inhibitors showed a reduction in mortality. Two patients were commenced on an ACE inhibitor and three had the dose increased during the three-month follow-up. Twenty-seven patients died during hospital admission and 22 during the three-month follow-up (40 over the age of 75 years).

Seventy-six patients were receiving antiarrhythmic treatment on discharge for atrial fibrillation (digoxin 61, amiodarone 5). Of these, 30 (39.8%) were taking anticoagulation to prevent embolic events. The notes in nine patients showed a contraindication for anticoagulation: gastrointestinal blood loss (4), malignancy (3) and other (2). The remaining 37 patients had no recorded contraindication but the majority were elderly (36>75 years and 21>80 years of age). All these patients were treated with aspirin instead.

Discussion

Our results clearly show that, on discharge from hospital, approximately one in four patients with heart failure and without a documented contraindication, are not prescribed an ACE inhibitor. This state of affairs is unlikely to be unique to this hospital. This is surprising in view of the overwhelming evidence that ACE inhibitors reduce fatal and nonfatal events in patients with overt heart failure and left ventricular dilatation in patients with asymptomatic left ventricular systolic dysfunction or following myocardial infarction.

The prevalence of heart failure continues to increase, in spite of better treatment of hypertension and other risk factors for coronary heart disease. ACE inhibitors have been shown to reduce or reverse the extent of ventricular dilatation and the incidence of reinfarction. Little is known, however, of the optimum dose of ACE inhibitor required to observe the expected improvement in cardiovascular events. Continuous inhibition of the angiotensin-converting enzyme is probably desirable. However, many of the patients who were prescribed an ACE inhibitor were given it at an inappropriate dose; only 16.7% of patients were on the doses used in the clinical trials showing a reduction in morbidity and mortality: CONSENSUS5 (10-20 mg enalapril twice a day); SOLVD6 (10 mg enalapril twice a day), SAVE7 (captopril 50 mg three times a day), and AIRE8 (5 mg ramipril twice a day). Similar results were also reported in 157 patients seen in tertiary referral clinics for heart failure. Even though no information is provided on the way the patients were selected, a wide range of different ACE inhibitor regimes was used, and only 46% of patients were receiving doses suggested by the results of survival trials.

The incidence of heart failure increases exponentially with age. It affects about 1% of people in their 50s and rises to 10% of those in their 80s. Therefore, to have an impact on the overall mortality and morbidity from heart failure one should aim to treat this latter group of patients. However, the published clinical studies have, to a large extent, excluded older patients: in the CONSENSUS study, the mean age was 71 years, in the SOLVD study the mean age was 61 years (patients over 80 were excluded), in the SAVE study the mean age was 59 years and in the AIRE study the mean age was 65 ± 11 years. There is, of
course, no reason to believe that elderly patients with heart failure are different from younger patients. Indeed, in the GISSI-3 trial, there was an even greater reduction in cardiovascular events in patients over 70 years, when compared to those under 70 years, who were treated with lisinopril following acute myocardial infarction.

Our results also suggest that coronary heart disease is the most frequently encountered cause of congestive heart failure in patients presenting to hospital. Furthermore, hypertension was found to be a predisposing risk factor in 25% of patients with heart failure. In the past, hypertension was the major contributor to the increased incidence of heart failure in the general population (preceding heart failure in approximately 75% of all cases). Although conclusive evidence is lacking, it has been postulated that the main reason to account for this change in the aetiology of heart failure, is the better management of severe hypertension over the last 30 years. Recent trials have also shown that treatment with warfarin significantly reduces the incidence of stroke among patients with atrial fibrillation (20–30 strokes prevented per 1000 patient years of treatment). As a result, there have been calls for the increased use of anticoagulant drugs in these patients. Even though our survey did not primarily examine the use of anticoagulation in patients with atrial fibrillation, it is clear that, at least in those patients with heart failure and atrial fibrillation, warfarin has been prescribed in the majority of patients to reduce the risk of embolic stroke; all but one patient not receiving warfarin was over 75 years of age where the adverse effects of anticoagulants are increased. However, it is increasingly recognised that warfarin is safe in selected elderly patients, and indeed the elderly may be the group that benefit most in terms of stroke prevention.

In conclusion, our results clearly demonstrate that in patients admitted to hospital with heart failure, one in four patients who should be receiving an ACE inhibitor by the time of discharge, are not. This is surprising in view of the overwhelming evidence that ACE inhibitors improve outcome in all patients with symptomatic or asymptomatic left ventricular dysfunction. Many of the patients who were prescribed an ACE inhibitor were given it at an inappropriate dose; only 24 patients (16.7%) were on the dose that was used in the clinical trials where ACE inhibitors showed a reduction in mortality. The average age of the patients in our study was 76 years, ie, an older age group than most of published studies with ACE inhibitors. Specific studies looking at the effect of ACE inhibitors in elderly patients would be helpful, as well as studies to determine the optimum treatment regimen for this age group.