Amlodipine or lisinopril was not better than chlorthalidone in lowering CHD risk in hypertension


**Question**
In predominately older patients with hypertension and ≥1 other coronary heart disease (CHD) risk factor, is amlodipine or lisinopril better than chlorthalidone in lowering the risk for CHD or other cardiovascular disease (CVD) events?

**Design**
Randomized (allocation concealed*), blinded [patients, clinicians, data collectors, outcome assessors, and steering committee]†,* controlled trial with mean 4.9-year follow-up.

**Setting**
623 centers in the United States, Canada, Puerto Rico, and the U.S. Virgin Islands.

**Patients**
33 357 patients ≥55 years of age (mean age 67 y; 53% men) who had stage 1 or stage 2 hypertension and ≥1 additional risk factor for CHD. The risk factors were myocardial infarction (MI) or stroke in the previous 6 months, left ventricular hypertrophy, type 2 diabetes, smoking, high-density lipoprotein cholesterol < 0.91 mmol/L, or other athero-sclerotic CVD. Patients with a history of heart failure (HF) or known left ventricular ejection fraction < 35% were excluded. Follow-up was 97%; all randomized patients were included in the analysis.

**Main Results**
Analysis was by intention to treat. Patients who received amlodipine or lisinopril did not differ from those who received chlorthalidone for the primary outcome (Table). The amlodipine group had a 38% (95% CI 25 to 52) relative risk increase for HF and a 35% (CI 21 to 50) greater risk for hospitalized fatal HF than did the chlorthalidone group. The lisinopril group had a 15% (CI 2 to 30) greater risk for stroke and a 10% (CI 5 to 16) greater risk for combined CVD than did the chlorthalidone group.

**Conclusions**
In older patients with hypertension and ≥1 other coronary heart disease (CHD) risk factor, amlodipine or lisinopril was not better than chlorthalidone in lowering the risk for CHD or other cardiovascular disease (CVD) events. Risks for some CVD events were greater with amlopidine or lisinopril.

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*See Glossary.
†Information provided by author.

### Amlodipine or lisinopril vs chlorthalidone for combined fatal CHD or nonfatal MI at mean 4.9 years‡

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Amlodipine</th>
<th>Lisinopril</th>
<th>Chlorthalidone</th>
<th>RRR (95% CI)</th>
<th>NNT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Amlodipine</td>
<td>11.3%</td>
<td>—</td>
<td>11.5%</td>
<td>2% (−7 to 10)</td>
<td>Not significant</td>
</tr>
<tr>
<td>Lisinopril</td>
<td>—</td>
<td>11.4%</td>
<td>11.5%</td>
<td>1% (−8 to 9)</td>
<td>Not significant</td>
</tr>
</tbody>
</table>

‡CHD = coronary heart disease; MI = myocardial infarction. Other abbreviations defined in Glossary.

**Commentary**
A study of astonishing methodologic beauty, ALLHAT is a large, simple, practice-based, double-blind, active-comparison trial that poses important public health questions about the effects of first-line antihypertensive therapies on major health outcomes, including MI, stroke, and HF. This combination of excellent methods and outstanding questions means that the findings merit careful attention, wide dissemination, and clinical translation.

In ALLHAT, neither the calcium-channel blocker amlodipine nor the ACE inhibitor lisinopril was better than the low-dose diuretic chlorthalidone for any primary or secondary outcome. Conversely, low-dose diuretics were superior to amlodipine or lisinopril for at least 1 major disease outcome. Low-dose diuretics alone should be the first-line drug therapy for hypertension.

For patients starting drug therapy, low-dose diuretics should be the initial therapy for uncomplicated hypertension, although it may take 4 weeks to achieve maximum effect. If blood pressure (BP) is not controlled on diuretics alone, a second drug should be added. Many patients are already treated, but not with a diuretic. For those with uncontrolled BP on a nondiuretic regimen, a low-dose diuretic should be added. If BP is controlled with a nondiuretic, the patient should be switched to a low-dose diuretic.

The ALLHAT trial, led by academic researchers and funded by the U.S. National Institutes of Health and Pfizer, represents an important university-industry collaboration: Newer drugs were compared with an established drug instead of a placebo for clinically important outcomes. Diuretics should be the standard reference group in future comparative trials.

The superiority of low-dose diuretics is not likely to be promoted by industry. Indeed, “opinion leaders” argue that BP control is inadequate; everyone needs 2 drugs, so combination therapy should be first-line therapy (1). During ALLHAT, which used the traditional stepped-care approach, 40% to 60% of patients were taking only 1 drug. For the large proportion of patients with hypertension whose BP can be controlled on 1 drug, the 2-drugs-for-everyone approach, which remains largely untested, would unnecessarily expose many patients to adverse effects without known benefit.

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**Reference**