Losartan reduced cardiovascular morbidity and mortality more than atenolol in patients with diabetes and essential hypertension


QUESTION: In patients with diabetes, essential hypertension, and signs of left ventricular hypertrophy (LVH), is losartan-based therapy more effective than atenolol-based therapy?

Design
Randomised (unclear allocation concealment*), blinded (patients and monitoring committee), controlled trial with ≥ 4 years follow up.

Setting
Multicentre trial in Europe and the United States.

Patients
1195 patients (a predefined subgroup of patients who had diabetes mellitus at the start of the LIFE study) who were 55 to 80 years of age (mean age 67 y, 53% women) with hypertension (sitting blood pressure [BP] after 1 to 2 wks of placebo of 160 to 200 mm Hg systolic, 95 to 115 mm Hg diastolic, or both) and electrocardiographic signs of LVH. Exclusion criteria included secondary hypertension; myocardial infarction (MI) or stroke within the previous 6 months; angina pectoris requiring treatment with β-blockers or calcium antagonists; and heart failure or left ventricular ejection fraction ≤ 40%. Follow up was 100%.

Intervention
Patients were allocated to losartan based therapy (n = 586) or atenolol based therapy (n = 609). Losartan and atenolol were started at 50 mg/day, combined with low dose hydrochlorothiazide if needed then increased to 100 mg/day if needed, and supplemented with other antihypertensives (except β-blockers, angiotensin-converting enzyme [ACE] inhibitors, or angiotensin-receptor blockers [ARBs]) to reach a target BP < 140/90 mm Hg.

Main outcome measures
The primary end point was a composite of cardiovascular mortality, MI, and stroke. One of the secondary end points was heart failure.

Main results
Analysis was by intention to treat. The composite end point, cardiovascular mortality, all cause mortality, and heart failure, occurred less frequently in patients assigned to losartan than in those assigned to atenolol.

Conclusion
In patients with diabetes, essential hypertension, and signs of left ventricular hypertrophy, losartan reduced cardiovascular morbidity and mortality and all-cause mortality more than atenolol.

*ASee glossary.

Losartan-based therapy v atenolol-based therapy in diabetes with essential hypertension and signs of left ventricular hypertrophy

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Losartan</th>
<th>Atenolol</th>
<th>RRR (95% CI)</th>
<th>NNT (CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Composite end point‡</td>
<td>18%</td>
<td>23%</td>
<td>22% (2 to 39)</td>
<td>21 (12 to 256)</td>
</tr>
<tr>
<td>Cardiovascular mortality</td>
<td>6%</td>
<td>10%</td>
<td>36% (5 to 57)</td>
<td>28 (18 to 211)</td>
</tr>
<tr>
<td>All-cause mortality</td>
<td>11%</td>
<td>17%</td>
<td>37% (15 to 53)</td>
<td>16 (12 to 40)</td>
</tr>
<tr>
<td>Heart failure</td>
<td>5%</td>
<td>9%</td>
<td>40% (8 to 61)</td>
<td>28 (19 to 145)</td>
</tr>
</tbody>
</table>

†Abbreviations defined in glossary; RRR, NNT, and CI calculated from data in article. ‡Cardiovascular mortality, stroke, and myocardial infarction.

COMMENTARY
Physicians can become frustrated when prescribing medications for asymptomatic patients with chronic diseases. They may doubt that the benefit of treatment exceeds the cost and risks for side effects. Patients may not comply if the cost or side effects exceed perceived benefits. As a result, therapeutic goals may be difficult to achieve.

The LIFE studies show that potentially life-threatening complications can be reduced with fewer side effects. The original, placebo controlled public health trials of hypertension used thiazides and β-blockers. These agents successfully reduced cardiovascular disease and stroke; thus, they have been considered first-line treatment for hypertension.1 In the LIFE studies, losartan reduced stroke and combined cardiovascular end points to a greater degree than atenolol. The data indicate that cardiovascular protection by using an ARB is superior to that of a β-blocker for patients with hypertension and LVH, which are independent risk factors for cardiovascular disease. Other trials using ACE inhibitors and ARBs have shown cardiovascular and renal benefits unrelated to their effects on hypertension.2–3 We need to reconsider the selection of first-line hypertensive therapy, particularly in patients at high risk.

Several comments should be made concerning the study methodology and results. Most patients required ≥ 2 agents to reach target level BP, which is consistent with other trials. Use of multiple agents could confound the comparison between the 2 agents in reducing risk. Lower systolic pressures should have been targeted. The final mean systolic BP was 146 mm Hg in patients with diabetes and 144 mm Hg in those without. To minimise cardiovascular events, systolic pressure should be decreased to 120 mm Hg and 140 mm Hg in patients with and without diabetes, respectively.1 16% of the patients smoked. No attempt to change dietary, exercise, or smoking habits was noted. Visits were semianual; more frequent appointments could produce better compliance, lower BP, and more positive lifestyle changes. Aggressive anti-hypertensive therapy, daily aspirin administration, and lifestyle improvements could have further reduced cardiovascular events.

Continued on next page
Losartan reduced strokes and new onset diabetes more than atenolol in essential hypertension


QUESTION: In patients with essential hypertension and signs of left ventricular hypertrophy (LVH), is losartan-based therapy more effective than atenolol-based therapy?

Design
Randomised (unclear allocation concealment*), blinded (patients and monitoring committee)*, controlled trial with ≥1 years follow-up.

Setting
Multicentre trial in Europe and the United States.

Patients
9229 patients 55 to 80 years of age (mean age 67 y, 54% women) with hypertension (sitting blood pressure [BP] after 1 to 2 wks of placebo of 160 to 200 mm Hg systolic, 95 to 115 mm Hg diastolic, or both) and electrocardiographical signs of LVH. Exclusion criteria included secondary hypertension; myocardial infarction (MI) or stroke within the previous 6 months; angina pectoris requiring treatment with β-blockers or calcium antagonists; and heart failure or left ventricular ejection fraction ≤40%. Follow-up was 99%.

Intervention
Patients were allocated to losartan based therapy (n = 4605) or atenolol based therapy (n = 4588). Losartan and atenolol were started at 50 mg/day, combined with low dose hydrochlorothiazide if needed and then increased to 100 mg/day if needed, and supplemented with other antihypertensive drugs (except β-blockers, angiotensin-converting enzyme [ACE] inhibitors, or angiotensin-receptor blockers [ARBs]) to reach a target BP <140/90 mm Hg.

Main outcome measures
The primary end point was a composite of cardiovascular mortality, MI, and stroke. One of the secondary end points was new onset diabetes.

Main results
Analysis was by intention to treat. The composite end point, fatal or non-fatal stroke and new onset diabetes, occurred less frequently in patients assigned to losartan than in those assigned to atenolol (table). No difference existed between the groups for cardiovascular mortality or MI. BP control, dose titration, and use of other antihypertensives were similar in both groups.

Conclusion
In patients with essential hypertension and signs of left ventricular hypertrophy, losartan reduced strokes and new-onset diabetes more than atenolol.

*See glossary.
Losartan reduced strokes and new onset diabetes more than atenolol in essential hypertension

Evid Based Med 2002 7: 174
doi: 10.1136/ebm.7.6.174

Updated information and services can be found at:
http://ebm.bmj.com/content/7/6/174

These include:

References
This article cites 6 articles, 1 of which you can access for free at:
http://ebm.bmj.com/content/7/6/174#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 (403)
- Stroke (546)
- Drugs: cardiovascular system (754)
- Epidemiologic studies (1092)
- Diabetes (365)
- Clinical trials (epidemiology) (1594)
- Ischaemic heart disease (419)
- Chemotherapy (82)
- Immunology (including allergy) (571)

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/