Combination ACE inhibitor and angiotensin receptor blocker therapy was better than monotherapy in non-diabetic renal disease


QUESTION: In patients with non-diabetic renal disease, what is the effectiveness of the angiotensin II receptor blocker (ARB) losartan, the angiotensin converting enzyme (ACE) inhibitor trandolapril, or the 2 drugs combined for delaying disease progression?

Design
Randomised (unclear allocation concealment†), blinded (clinicians, patients, data collectors, and monitoring committee),† controlled trial with 3 years of follow up.

Setting
Hospital outpatient renal clinic serving 3 cities in Japan.

Patients
301 patients 18–70 years of age who had chronic non-diabetic renal insufficiency, persistent proteinuria, and no history of allergic reaction to drugs. Exclusion criteria included immediate need for renal replacement therapy; need for corticosteroids, non-steroidal anti-inflammatory drugs, or immunosuppressive drugs; proteinuria >10 g/day and hypoalbuminaemia <28 g/l; other serious disease; and pregnancy or breastfeeding. 263 patients (mean age 45 y, 54% men) completed an 18 week run in period and were randomised. Follow up of these patients was 97%.

Intervention
Patients were allocated to losartan, 100 mg/day plus placebo (n=89); trandolapril, 3 mg/day, plus placebo (n=86); or a combination of losartan, 100 mg, and trandolapril, 3 mg/day (n=88).

Main outcome measures
A combined endpoint of time to doubling of serum creatinine concentration or end stage renal disease (ESRD) (glomerular filtration rate <7 ml/min per 1.73 m² or implementation of dialysis). Secondary outcomes were changes in blood pressure and urinary protein excretion, and adverse effects.

Main results
Analysis was by intention to treat. At 3 years, fewer patients who received combination treatment reached the combined endpoint than did patients who received either drug with placebo (table). Blood pressure did not differ between groups. Patients in the combination treatment group had the greatest decrease in urinary protein excretion rate (maximum decrease 75%) compared with losartan alone (42%) and trandolapril alone (44%). Groups did not differ for adverse effects, and no patient had an acute decline in renal function.

Conclusion
In patients with non-diabetic renal disease, losartan and trandolapril combined were better than either drug alone for delaying disease progression.

*See glossary.

Combination treatment vs losartan or trandolapril alone for non-diabetic renal disease at 3 years

<table>
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<tr>
<th>Outcome</th>
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<td>Combined endpoint</td>
<td>Combination treatment vs losartan</td>
<td>11% vs 23%</td>
<td>49% (0.5 to 75)</td>
<td>9 (5 to 1722)</td>
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<tr>
<td>Combination treatment</td>
<td>v trandolapril</td>
<td>11% vs 23%</td>
<td>50% (1.7 to 75)</td>
<td>9 (5 to 369)</td>
</tr>
</tbody>
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†Combined endpoint = time to doubling of serum creatinine concentration or end stage renal disease. Abbreviations defined in glossary; RRR, NNT, and CI calculated from data in article.

COMMENTARY
ACE inhibitors are standard treatment for slowing the progression of non-diabetic renal disease and this benefit is mediated by factors in addition to lowering blood pressure and urinary protein excretion.1 The possibility that additional reductions in the rate of progression are achievable has led to consideration of the combination of an ACE inhibitor with an ARB. Several small short term studies have shown an improved antiproteinuric effect and the safety of this combination treatment.

The COOPERATE study confirms these findings and reports a remarkable reduction of the long term combined outcome of doubling of serum creatinine or ESRD with combination treatment compared with either drug alone. Only 1 of 85 patients in the combination treatment group reached ESRD at 3 years. These results were achieved with excellent blood pressure control (125/72 mm Hg during the trial), marked dietary protein restriction, and the use of maximum doses of both classes of drugs.

Previous studies of ACE inhibitors2–3 were designed to achieve diastolic blood pressures <90 mm Hg with submaximal doses and showed less compliance with low protein diets than did the COOPERATE trial. The generalisability of this study is hampered by the relatively young age of the patients (mean age 45 y) and the preponderance of glomerular renal disease, primarily IgA nephropathy, an uncommon cause of ESRD in Europe and North America. Future studies evaluating combination treatment need to include older patients with underlying nephrosclerosis before this promising approach can be recommended as a safe and effective therapy for non-diabetic renal disease.

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