

COMMENTS AND RESPONSES

Comment on: Hellemons et al. Initial Angiotensin Receptor Blockade- Induced Decrease in Albuminuria Is Associated With Long-Term Renal Outcome in Type 2 Diabetic Patients With Microalbuminuria: A Post Hoc Analysis of the IRMA-2 Trial. Diabetes Care 2011; 34:2078-2083

Hellemons et al. (1) in a post hoc analysis of the Irbesartan in Patients with Type 2 Diabetes and Microalbuminuria (IRMA)-2 trial provocatively showed that the initial reduction of microalbuminuria with an angiotensin receptor blocker (ARB) was independently

associated with renoprotection (i.e., independent of blood pressure changes). Anti-hypertensive medication was removed for a 3-week run-in period before the ARB was (re)introduced. If these results could be confirmed in a randomized controlled trial (RCT) it would imply that aggressive reduction of microalbuminuria should be attempted—an approach not currently recommended by the American Diabetes Association (2). This might be a difficult goal to accomplish clinically given the marked day-to-day intraindividual variability (33–61%) of microalbuminuria (3,4). We could not lower established microalbuminuria in an RCT pilot study in patients already treated with submaximal doses of an ACE inhibitor by maximizing the doses of a combination of benazepril plus losartan compared with 10 mg of the ACE inhibitor over a mean of 12 months (5). In our run-in period, patients were kept on 10 mg of benazepril. This might be an important factor in designing future real-world RCTs to test the hypothesis generated by the post hoc analysis of the IRMA-2 trial (1).

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