Inhibition of RAS in diabetic nephropathy

Rabi Yacoub  
Kirk N Campbell

Department of Medicine, Division of Nephrology, Icahn School of Medicine at Mount Sinai, New York, NY, USA

Abstract: Diabetic kidney disease (DKD) is a progressive proteinuric renal disorder in patients with type 1 or type 2 diabetes mellitus. It is a common cause of end-stage kidney disease worldwide, particularly in developed countries. Therapeutic targeting of the renin–angiotensin system (RAS) is the most validated clinical strategy for slowing disease progression. DKD is paradoxically a low systematic renin state with an increased intrarenal RAS activity implicated in its pathogenesis. Angiotensin II (AngII), the main peptide of RAS, is not only a vasoactive peptide but functions as a growth factor, activating interstitial fibroblasts and mesangial and tubular cells, while promoting the synthesis of extracellular matrix proteins. AngII also promotes podocyte injury through increased calcium influx and the generation of reactive oxygen species. Blockade of the RAS using either angiotensin converting enzyme inhibitors, or angiotensin receptor blockers can attenuate progressive glomerulosclerosis in animal models, and slows disease progression in humans with DKD. In this review, we summarize the role of intrarenal RAS activation in the pathogenesis and progression of DKD and the rationale for RAS inhibition in this population.

Keywords: renin–angiotensin system, diabetic kidney disease, angiotensin II, angiotensin-converting enzyme inhibitors, angiotensin receptor blockers

Introduction

Worldwide, more than 347 million people are suffering from diabetes mellitus (DM) and its complications which are projected to be the seventh leading cause of death by 2030. The incidence has doubled in the last two decades, and DM is the underlying cause of both micro- and macrovascular disorders such as diabetic kidney disease (DKD), retinopathy, coronary artery disease, and peripheral vascular disease. DKD is a progressive disease that affects patients with long-standing type 1 (insulin dependent) or type 2 (non-insulin-dependent) DM, and is functionally characterized by different degrees of albuminuria and chronic kidney disease. Associated histological structural changes include mesangial expansion, glomerular basement membrane thickening, glomerular sclerosis, and in advanced stages, tubulointerstitial fibrosis. In the US, it is estimated that 44% of all new end-stage renal disease (ESRD) cases are due to DKD. Interestingly, despite the increased incidence rate of DM, recent studies have shown that the incidence of DKD has dramatically decreased with the implementation of tight glycemic control and the use of renin–angiotensin system (RAS) blockers. The use of RAS blockers in DKD not only decreases albuminuria, but slows disease progression, which is only partially explained by their hemodynamic effects. The RAS cascade starts with the production of prorenin in the juxtaglomerular cells, which...
is cleaved into renin.\(^8\) Renin then cleaves angiotensinogen (AGT) to form angiotensin I (AngI) which later is converted by angiotensin converting enzyme (ACE) to the octapeptide angiotensin II (AngII). AngII in turn activates both angiotensin type 1 and 2 (AT\(_1\), AT\(_2\), respectively) receptors (Figure 1). AngII was previously considered to be the only active metabolite of this cascade, but evidence in the last few decades has shed light on the importance and biological role of each product including prorenin, renin, and AngI. In this review, we summarize the role of RAS in diabetic nephropathy and the mechanisms by which RAS inhibition slows disease progression.

**Genetics of RAS**

Host genetic variation in RAS components may predict the risk of developing DKD. It is well documented that there is an increased likelihood of developing DKD in family members who have siblings affected with the disease. Genetic studies have sought to determine variants and factors predisposing individuals to DKD.\(^9–11\) Some studies examined the role of the ACE gene genotype as a potential genetic risk factor and showed conflicting results. Investigators analyzed the insertion/deletion (I/D) and DD polymorphism in ACE gene in type 2 DM patients and concluded that DD polymorphisms are associated with greater risk of renal function decline, severe proteinuria, and worse outcomes.\(^12–14\) These findings were not seen in type 1 DM patients or in a large review of 19 studies that included type 1 and 2 patients.\(^15–17\) Male, and not female, patients with type 1 diabetes and the AA haplotype of the AT\(_2\) receptor gene on the X-chromosome had a lower glomerular filtration rate than those with the GT haplotype.\(^18\)

**Hemodynamic effects of RAS on the glomerular filter**

Renal autoregulation enables the kidneys to maintain a stable intraglomerular pressure and glomerular filtration rate (GFR) as the mean arterial pressure varies from 80–160 mmHg.\(^19\) This is achieved through tubular–glomerular feedback (TGF) and myogenic response affecting the afferent arteriole’s tone, and angiotensin II (AngII) affecting the tone of both the afferent and the efferent arterioles.\(^20\) This autoregulatory mechanism is disrupted in different diseases, including hypertension, chronic kidney disease, and diabetes (Figure 2). Early in the course of DKD, an elevation in the GFR is seen, accompanied later by glomerular and renal hypertrophy along with increased intraglomerular pressure. This in turn increases the shear stress on the glomerular capillary wall resulting in progressive and sustained renal damage.\(^21\)

A key mechanistic factor in this hyperfiltration response in DKD is RAS mediated glomerular afferent arteriole dilatation which increases renal blood flow and intraglomerular pressure.\(^22\) Other contributors to increased afferent arteriole dilatation include insulin-like growth factor I (IGF-1),\(^23\) atrial natriuretic peptide,\(^24\) and advanced glycation end products, the latter also influencing the micro- and macrovascular changes seen in DM.\(^25\) Additionally in DM, there is an augmented sodium–glucose cotransport leading to volume expansion, along with hyperinsulinemia and local AngII stimulating sodium reabsorption in the proximal tubule.\(^26–28\) This increased proximal sodium reabsorption activates the tubuloglomerular feedback mechanism in the macula densa, and can then raise the GFR via dilatation of the afferent arteriole.\(^26\)

AngII has vasoconstrictive effects on the afferent and the efferent arterioles as well as the interlobular arteries, mediated in part by the local generation of thromboxane A2. The constrictive efficacy is greatest at the efferent arteriole due to its smaller basal diameter. This leads to a net reduced renal blood flow, increased intraglomerular pressure, and an increase in GFR.\(^29\) Concurrently, AngII stimulates the release of vasodilator prostaglandins from the glomeruli and constricts glomerular mesangial cells, thereby lowering the
surface area available for filtration. AngII also sensitizes the afferent arteriole to the constricting signal of tubuloglomerular feedback.30-33

**RAS activation and its importance in DKD**

The beneficial effects of RAS blockers in slowing the progression of diabetic nephropathy have been well documented,34-36 despite the low systemic renin state in DKD. This phenomenon is thought to be reflecting the activated intrarenal renin system or increased intrarenal sensitivity to AngII.37 Likely reflecting different stages of kidney disease and different measurement methods (renin activity, protein level, RNA expression, serum potassium and bicarbonate, and immunohistochemistry and fluorescence), human clinical data measuring RAS activation have been conflicting in DKD.38-41 Experimental models however tend to show a consistent increase in RAS activation.42-46 This activation in RAS involves all factors and steps in the cascade and takes place in a paracrine fashion. Table 1 summarizes the deleterious effects of RAS cascade activation in DKD.

**Prorenin**

The importance of prorenin and its biological role beyond the formation of renin was first established in the 1980s when it was found that its levels, despite the low systemic renin levels and activity, are increased in diabetic subjects, and correlate with the presence of retinopathy and nephropathy.39

**Table 1** The deleterious effects of RAS cascade activation in DKD

<table>
<thead>
<tr>
<th>RAS Cascade Component</th>
<th>Effect on DKD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Prorenin</td>
<td>Glomerulosclerosis</td>
</tr>
<tr>
<td></td>
<td>Renal fibrosis</td>
</tr>
<tr>
<td>Renin</td>
<td>Nonenzymatic enhancement of angiotensin II synthesis</td>
</tr>
<tr>
<td></td>
<td>Mesangial hypertrophy</td>
</tr>
<tr>
<td></td>
<td>Renal fibrosis</td>
</tr>
<tr>
<td>Angiotensin II</td>
<td>Increased intraglomerular pressure</td>
</tr>
<tr>
<td></td>
<td>Water and solute retention</td>
</tr>
<tr>
<td></td>
<td>Mesangial cell proliferation and hypertrophy</td>
</tr>
<tr>
<td></td>
<td>Renal fibrosis</td>
</tr>
<tr>
<td></td>
<td>Glomerular and tubulo-interstitial microinflammation</td>
</tr>
<tr>
<td></td>
<td>Insulin resistance</td>
</tr>
<tr>
<td></td>
<td>Decreased AT,R synthesis</td>
</tr>
</tbody>
</table>

**Abbreviations:** AT,R, angiotensin type 2 receptor; RAS, renin–angiotensin system; DKD, diabetic kidney disease.
In DM, prorenin represents 95% of circulating renin, and the mechanism by which prorenin is implicated in the development of DKD and retinopathy in this population is under investigation. It binds to the prorenin receptor (PRR) and, independently of AngII, triggers intracellular signaling and the activation of the mitogen-activated protein (MAP) kinases ERK1/2, leading to upregulation of TGFβ1, PAI1, collagens, fibronectin, and cyclooxygenase-2. This raises the possibility that high prorenin levels play a role in diabetic nephropathy by stimulating PRR and inducing profibrotic protein syntheses. Indeed, transgenic rats overexpressing human PRR develop proteinuria, and a slowly progressive glomerulosclerosis and kidney damage independent of AngII.

Interestingly, inductible overexpression of prorenin does not result in glomerulosclerosis suggesting that prorenin may augment the fibrosis rather than causing it. Prorenin also participates in the generation of AngI. Prorenin binding to PRR induces a conformational change involving unfolding of the peptide from the enzymatic cleft so the cleft is now accessible to AGT and generates AngI.

**ACE**

The best evidence for intrarenal renin production and activity in DM comes from spontaneous or streptozotocin (STZ)-induced diabetes animal models. In spite of the suppressed or normal plasma renin activity, the intrarenal Ren mRNA and protein levels, most significantly in the proximal tubule, are increased in DM. The total renal ACE activity is significantly reduced in DM rats, with specific redistribution in diabetic kidneys. While proximal tubule ACE activity is reduced, ACE staining intensity is enhanced in diabetic glomeruli and renal vasculature. This suggests a role for glomerular ACE in mediating nephron injury, possibly by increasing local intraglomerular AngII formation.

**Renin**

Early diabetes causes a significant stimulation of the proximal tubule renin mRNA expression. Renin, independent of its enzymatic action to enhance AngII synthesis, directly increases production of transforming growth factor β (TGF-β), the fibrogenic cytokine. Renin binds to its specific receptor on the cell surface of mesangial cells, leading to hypertrophy, and enhanced efficiency of angiotensinogen cleavage by renin, thereby unmasking prorenin catalytic activity. The renin receptor was also localized in the subendothelium of the renal arteries suggesting that renin has a novel receptor-mediated action that could play a role in renal fibrosis. In podocytes, high glucose is shown to cause increased AngII generation through increasing renin mRNA expression with a concomitant increase in PRR and hence augmenting the conversion from AGT to AngI.

**Angiotensinogen**

The high extracellular glucose in DM stimulates the synthesis of AGT in a concentration dependent manner and increases the expression of its gene (Agt) in the proximal tubule of the kidneys. This increased AGT protein level in the proximal tubules is due to many factors, including the effects of high extracellular glucose on Agt expression via reactive oxygen species and the direct effect of glucose on its promoter. A glucose response element has been located on the Agt promoter.

**Angiotensin II**

As previously mentioned, in diabetic nephropathy there is an increase in the generation of the intrarenal AngII despite the systemic suppression of RAS. The deleterious effects of this rise in AngII go beyond the hemodynamic changes to involve insulin resistance, growth promotion, and tubular damage. One of the most important roles of AngII in DKD is its association with volume expansion through water and Na reabsorption. It activates the Na+/H+ antiporter in the luminal membrane through stimulation of an inhibitory G protein that decreases cyclic AMP (adenosine monophosphate) generation, minimizing the normally suppressive effect of cyclic AMP on Na+/H+ exchange. AngII stimulates phosphatidylinositol turnover, resulting in the generation of protein kinase C. It also increases the secretion of aldosterone from the adrenal cortex enhancing Na+ transport in the cortical collecting tubule. AngII inhibits protease activity in the proximal tubule and causes mesangial cell expansion via decreasing the activity of plasminogen activator. AngII mediated TGF-β1 upregulation and vascular endothelial growth factor release from the glomerular epithelial and mesangial cells contribute to mesangial matrix expansion. Renal fibroblasts express AT1 receptor and respond to AngII stimuli by cell proliferation, matrix expansion, and synthesis of fibronectin, by a TGF-β-dependent mechanism. Microinflammation of the glomeruli and tubulointerstitial regions and subsequent extracellular matrix expansion are common pathways for the progression of DKD. AngII activates inflammatory cells by direct chemotaxis including osteopontin (OPN), RANTES and the production of other proinflammatory mediators, including MCP-1 and TGFβ. It activates protein kinase C, protein tyrosine kinases (PTK), mitogen-activating protein kinases (MAPK), extracellular...
signal-regulated kinase (ERK), c-Jun amino terminal kinase (JNK), p38 MAP kinase (p38 MAPK), and the activator protein-1 (AP-1). These factors are implicated and involved in proliferation, differentiation, fibrosis, and inflammation processes. AngII has also been implicated in insulin resistance. AngII inhibits insulin-mediated GLUT4 translocation in this skeletal muscle model through a transient activation of ERK1/2 inhibiting insulin receptor substrate 1/2 (IRS-1/2) and through a direct inhibitory nitration of Akt. It induces tyrosine phosphorylation of IRS-1 by Janus kinase 2 associated with AT1 receptor stimulation which attenuates insulin-induced activation of phosphatidylinositol-3-kinase associated with IRS-1, leading to decreased insulin sensitivity. Treatment with the angiotensin II receptor blocker (ARB) candesartan increases renal insulin receptor expression in STZ-induced diabetic rat models and the insulin resistant Zucker rats irrespective of insulin levels.

It appears that high glucose directly stimulates podocytes to increase AngII production independent of ACE activity, and increases AT1 receptor levels promoting the deleterious effects of AngII on podocytes. AngII directly promotes podocyte injury via release of calcium from intracellular stores as well as influx from the extracellular space. Finally, AngII activates enzyme systems like Nox4 that use NADPH oxidases as substrates for superoxide generation. These mechanisms contribute to proteinuria and progressive renal disease.

Aldosterone

Aldosterone secretion from the adrenal cortex is mainly mediated by AngII and plasma potassium concentration. This secretion might result in a vicious cycle of local RAS stimulation causing further RAS activation and aldosterone generation. The effects of aldosterone go beyond solute and water retention, vasoconstriction, and subsequently, increased blood pressure. In the diabetic kidney, aldosterone activity contributes to tubulointerstitial fibrosis and inflammation, key features of disease progression. Aldosterone increases extracellular matrix production through enhanced TGF-β1 and type IV collagen gene expression. In diabetic rats, aldosterone induces glomerular and tubulointerstitial macrophage infiltration through TGFβ. Aldosterone also stimulates ROS production through activation of NADPH oxidase in mesangial cells. Systemically, higher circulating levels of aldosterone are associated with insulin resistance. There is a need for large prospective trials to determine the efficacy of aldosterone inhibition in the treatment of DKD.

ACE2 and angiotensin 1–7 (Ang 1–7)

Angiotensin converting enzyme 2 (ACE2) is an enzyme related to ACE, expressed in the intrarenal endothelium and the tubular epithelial cell. It hydrolyzes AngI to Ang 1–9, which is subsequently cleaved by ACE into Ang 1–7. ACE2 also catalyzes the conversion of AngII to Ang 1–7 (the major pathway of Ang 1–7 production). ACE2 does not produce AngII, or metabolize bradykinin (BK), and is not inhibited by ACE inhibition. Cardiac ischemia studies suggest that ACE2 and ACE have counterbalancing effects. Animal models with disrupted ACE2 gene showed significant cardiac abnormalities including decreased contractility and increased damage, probably mediated by the upregulation of hypoxia-induced genes. These histological findings were abrogated by concomitant ACE gene ablation. Ang 1–7 causes intrarenal vasodilation and counterbalances the vasoconstrictor effects of AngII via stimulation of NO and prostaglandins, and synergistically acts with BK. The effects are not limited to hemodynamics. In animal models of DKD, Ang 1–7 treatment ameliorates mesangial expansion and renal fibrosis, and reduces oxidative stress, inflammation, and lipotoxicity through blocking the activation of MAPKs by AngII and selective inhibition of glucose-stimulated protein synthesis, TGF-β1/Smad3- and VEGF-mediated pathways. There is a dynamic reciprocal interaction between Ang 1–7 and ACE; Ang 1–7 inhibits ACE by 30%–70%, and at the same time is degraded by it. In addition to its vasodilatory and natriuretic actions, Ang (1–7) has an antiproliferative effect on vascular smooth muscle cells.

Types of RAS blockades and their pleiotropic effects

Direct renin inhibitors (DRI), angiotensin converting enzyme inhibitors (ACEi), and angiotensin II receptor blockers (ARB) are available to treat hypertension and proteinuric kidney diseases. The use of these medications results in either decreased AngII (DRI and ACEi) or blocking its receptor AT1 (ARB). To better understand the differences in effects and actions of these medications, attention should first be paid to AngII receptor distribution. AT1 receptors are distributed throughout the kidney compartments. They are localized in the afferent and efferent arterioles, vascular smooth muscle cells, podocytes, glomerular mesangial and macula densa cells, the proximal tubule cell brush border, basolateral membranes, thick ascending limb epithelia, distal tubules, and the cortical collecting ducts. The AT1 receptor is localized in the afferent arteriole, glomerular...
endothelial and mesangial cells, proximal tubule cells, and interstitial cells.\textsuperscript{81,82} AT\textsubscript{1} receptor stimulation leads to renal arterial vasoconstriction, tubule epithelial sodium reabsorption, augmentation of tubulo–glomerular feedback sensitivity, and inhibition of pressure-natriuresis.\textsuperscript{77} Conversely, AT\textsubscript{2} receptor activation exerts the opposite effects with respect to cardiovascular hemodynamics and cell growth. AT\textsubscript{2} receptor activation stimulates BK and NO production, vasodilatation, and modulates the vasoconstrictor action mediated by the AT\textsubscript{1} receptor through endothelium-dependent vasodilatation.\textsuperscript{83}

DRI
AngII has a negative feedback on RAS, and during ACEi therapy, there is a rise in plasma renin activity (PRA) and concentration (PRC) as a result of decreased renal parenchymal AngII. This led to the development of DRIs for monotherapy or in combination with ACEi therapy. They have been shown to have a similar antiproteinuric and antihypertensive effects, but without significant cardiovascular benefits.\textsuperscript{84,85}

ACEi
ACEi is probably one of the most studied and used class of antihypertensive medication. Since the late 1970s and the use of captopril, ACEi has become the cornerstone in treating hypertensive patients with proteinuria. It inhibits ACE activity by blocking the conversion from AngI to AngII, and inhibits BK degradation. Bradykinin, via its B2 receptors, stimulates NO, cGMP, prostaglandin E2, and prostacyclin. Therefore, ACEi therapy not only inhibits AngII production but also increases the production of the vasodilatory factors.\textsuperscript{86} ACEi therapy also inhibits the ACE mediated degradation of Ang 1–7 augmenting the renoprotective effects of Ang 1–7. The pleiotropic effects of ACEi include anti-inflammatory and antioxidant,\textsuperscript{87} antithrombotic, and profibrinolytic activities.\textsuperscript{88} ACEi is also important in significantly improving arterial compliance through cytoprotection of vascular endothelium,\textsuperscript{89} along with better regression of left ventricular hypertrophy compared to beta blockers.\textsuperscript{90}

ARB
The major clinical and physiological differences between ACEi and ARBs are related to activation of AT\textsubscript{1} receptors. ARBs selectively block AT\textsubscript{1} receptor without directly affecting the synthesis of AngII and subsequent AT\textsubscript{2} activation. Some of the beneficial effects of using ARB are thought to be due to selective inhibition of AT\textsubscript{1} and the concomitant renoprotective stimulation of AT\textsubscript{2} receptors.

It has been proposed that ARBs might have a greater potential in preventing renal interstitial fibrosis compared to ACEi mediated by AT\textsubscript{2} receptor proapoptotic properties.\textsuperscript{91} This is supported by the fact that renal interstitial fibrosis after ureteral obstruction is enhanced in rats with either AT\textsubscript{2} deletion, mutation, or pharmacologic blockade.\textsuperscript{92,93}

RAS blockers and renal outcome studies
The effects and roles of RAS blockers in the progression of kidney disease are probably mostly seen due to their hemodynamic effects and blood pressure control. It is well known that blood pressure control is considered a cornerstone in the management and prevention of chronic kidney diseases.\textsuperscript{94} Thus, different studies have evaluated the hypothesis whether a stricter lower blood pressure control will confer a better renal outcome, and initial societies guidelines\textsuperscript{95–97} have recommended a lower blood pressure target (<130 mmHg systolic) for patients with CKD and significant proteinuria, based on the findings of post hoc analysis of major trials like MDRD\textsuperscript{98–100} AASK,\textsuperscript{101,102} and a patient level meta-analysis of eleven randomized trials.\textsuperscript{103} These studies mostly included nondiabetic kidney disease patients, and thus it is hard to extrapolate their findings and assume that DKD patients will respond similarly. To address this issue, post hoc analysis of RENAAAL\textsuperscript{104} and Irbesartan Diabetic Nephropathy Trial (IDNT)\textsuperscript{105} found that higher BP targets (140–149 mmHg for RENAAAL, and >145 mmHg for IDNT) resulted in higher risk of composite outcome. These studies also found that a tighter BP control (systolic <120 mmHg) resulted in increased mortality risk. It is important to mention that these studies were post hoc analyses that included a RAS blockers arm and a placebo arm, and were not designed to answer the question whether achieving a lower BP goal (<130 mmHg systolic) by increasing RAS blockers dose will have any benefit in this population. At the same time, these data were later challenged and questioned by the findings of increased risk of adverse outcomes (cardiac events, stroke, and falls) in different populations (especially elderly) and current guidelines have raised the target to 140 mmHg systolic.\textsuperscript{106–109}

Several studies have evaluated the role of ACEi in halting DKD progression. The Captopril-Diabetes study evaluated type 1 diabetic patients with nephrotic range proteinuria and showed a significant reduction in proteinuria along with lower GFR decline rate compared to placebo.\textsuperscript{110,111} The ADVANCE trial compared perindopril–indapamide combination to placebo in over 11,000 patients with DM, and found a significantly lower microalbuminuria, and lower worsening
The importance and significance of the effects of proteinuria reduction on renal outcomes was questioned since the treatment groups had better blood pressure control, which might have played a big role in preventing disease progression. Later studies showed that lowering proteinuria in patients with diabetic nephropathy (regardless of the agent used) is associated with improved outcomes and a decreased risk of ESRD.\(^{114\,\text{-}\,117}\)

The IDNT and the RENAAL trial studied ARBs therapy in patients with DKD and found similar beneficiary results. ARB therapy was superior to placebo and amlodipine in decreasing proteinuria, lower rates of doubling creatinine, or the development of ESRD and mortality.\(^{34\,\text{-}\,118}\)

No significant difference was found between ACEi and ARB in patients with early DKD. In the DETAIL trial, both groups (enalapril vs telmisartan) had similar 5-year GFR decline, annual changes in the GFR, blood pressure, serum creatinine, urinary albumin excretion, ESRD, cardiovascular events, and mortality.\(^{119}\)

**Trials evaluating dual RAS blockades and controversies**

It was previously demonstrated that ACEi/ARB combination therapy decreases proteinuria, more than monotherapy with either drug.\(^{120\,\text{-}\,122}\) These studies found no difference in adverse events in the combination group compared to monotherapy, but bore two major limitations that made their results and conclusion debatable. The sample sizes in these studies were small and patients were followed for a maximum of 16 weeks. Large scale randomized controlled trials conducted in different patient populations over a longer follow-up period failed to show any benefit on hard outcomes, including 50% drop in GFR, ESRD, and mortality.\(^{123\,\text{-}\,124}\) Patients on dual therapy had significantly higher adverse events, including hyperkalemia and hospitalization for acute kidney injury (AKI), as seen in the VA NEPHRON-D study,\(^{123}\) and even a statistically nonsignificant trend of higher incidence of chronic dialysis, doubling in serum creatinine and death, along with a statistically significant higher incidence rate of AKI requiring dialysis, hypotension, and hyperkalemia (ONTARGET trial).\(^{124}\) The results from studies evaluating the benefit of adding DRI (Aliskiren) to either ACEi or ARB in patients with DKD showed similar trend.\(^{125\,\text{-}\,126}\) Initially, the AVOID trial studied the combination between DRI and ARB in a group of 599 patients for 6 months and concluded that aliskiren may have renoprotective effects that are independent of its blood pressure-lowering effect in patients with hypertension and DKD, without a significant increase in adverse events.\(^{125}\) Later, the ALTITUDE investigators evaluated 8,561 patients for a median follow-up of 32.9 months and found that the combination therapy increased the risk of composite primary endpoint (ESRD, doubling of serum creatinine, renal death, cardiovascular death, cardiac arrest, heart failure, nonfatal myocardial infarction, or non-fatal stroke), along with hyperkalemia.\(^{126}\)

As many as 40% of subjects treated with RAS blockers (ACEi or ARBs) have above normal limits of aldosterone (aldosterone breakthrough).\(^{127\,\text{-}\,128}\) This is thought to be due to different mechanisms that include: stimulation from tissue not inhibited by ACEi/ARBs, AngII generation via pathway not requiring ACE, or other stimuli such as high serum potassium concentrations.\(^{129}\) Adding spironolactone (aldosterone receptor antagonist) to ACEi in DN patients with aldosterone breakthrough can decrease proteinuria.\(^{128}\) This study, though promising and reporting no significant change in serum potassium, was limited to a small number of patients and was of short duration. Similar findings on proteinuria were noted in different small and short duration randomized controlled studies using a combination therapy of spironolactone and either ACEi or ARBs.\(^{130\,\text{-}\,137}\) Not surprisingly, all studies reported either a trend or a statistically significant higher levels of serum potassium, and greater eGFR decline in the combination therapy group. Similar findings were noted in adding eplerenone (aldosterone antagonist) to ACEi or ARBs in patients with DN and preserved renal function.\(^{136}\)

There are no long-term data regarding the beneficial effects of adding aldosterone antagonist, and the current literature suggests possible increased risk of adverse events.

**Adverse events**

The major adverse events in patients using RAS are hypotension, hyperkalemia, angioedema, and reduction in GFR. ACEi can result in dry cough in up to 20% of cases, while ARB is associated with cough in up to 9.9% of cases.\(^{137\,\text{-}\,138}\) Recently there is an emerging concern about the development of sprue-like enteropathy in patients using olmesartan.\(^{139}\) Patients with bilateral renal artery stenosis, polycystic kidney disease, and advanced kidney disease may sustain a significant decrease in GFR, and in some cases AKI. The intrarenal perfusion pressure in these cases is already reduced, and patients are highly dependent on AngII to maintain a stable intraglomerular pressure and GFR. Similar but modest intrarenal hemodynamic changes are seen in elderly and DKD patients on RAS, increasing their risk of sustaining AKI in the setting...
of volume depletion, contrast agents, and major stressors like surgery.  

Summary and recommendations

Intrarenal activation of RAS in DKD plays a major role in disease pathophysiology and is implicated in its progression. The intrarenal generation is increased despite the systemic RAS suppression. This activation further downregulates the expression of both AT₁ and AT₂ receptors contributing to early diabetic nephropathy. This reduces AT₂ receptor mediated beneficial counterregulatory actions. The negative impact of increased AngII on the kidneys goes beyond the regulation of intraglomerular pressure. It leads to increased insulin resistance, solute retention, stimulation of TGF-β secretion, increased synthesis, and decreased degradation of matrix proteins, all leading to matrix accumulation, kidney fibrosis, and further kidney damage. The use of ACEi and ARBs has significantly changed the incidence of ESRD and has provided renal protection through their effects on RAS, hemodynamic changes, and other pleiotropic effects that include anti-inflammatory, antioxidant, antithrombotic effects, and profibrinolytic activities. Even though a combination therapy of both ACEi and ARBs has decreased protein excretion in DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.

Studies to date have found no difference in renal outcomes between ACEi and ARBs, thus all hypertension patients with proteinuric kidney disease should be started on either ACEi or ARBs to halt disease progression. In our opinion, we believe that the evidence and the risk of adverse events outweigh the benefits of RAS blockers combination therapy. This therapy, though resulting in decreased proteinuria, carries a significant risk of adverse outcomes including hyperkalemia, increased hospitalization events, acute kidney injury, and a faster decline in renal function, and should be avoided. Another important issue we as clinicians face on a daily basis is the increase in serum potassium level in patients on RAS blockers. Little is available in the literature in regards to the use of sorbitol-free DKD patients, it is associated with worse renal outcomes compared to monotherapy. Studies evaluating other components of the RAS, including ACE2 and Ang 1–7 along with genetic animal models are warranted.


The International Journal of Nephrology and Renovascular Disease is an international, peer-reviewed open-access journal focusing on the pathophysiology of the kidney and vascular supply. Epidemiology, screening, diagnosis, and treatment interventions are covered as well as basic science, biochemical and immunological studies. The journal welcomes original research, clinical studies, reviews & evaluations, expert opinion and commentary, case reports and extended reports. The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit http://www.dovepress.com/testimonials.php to read real quotes from published authors.

Submit your manuscript here: http://www.dovepress.com/international-journal-of-nephrology-and-renovascular-disease-journal