Differential Regulation of Renal Angiotensin Subtype AT1A and AT2 Receptor Protein in Rats With Angiotensin-Dependent Hypertension

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Abstract—This study was designed to investigate distribution and regulation of the renal AT1A and AT2 subtype receptors in rats with either systemic angiotensin II (Ang II)–induced hypertension or acute phase renal hypertension (2-kidney, 1-clip [2K1C] or 2-kidney, 1-figure-of-8-wrap [2K1W]). In normal rat kidneys, positive immunostaining for the AT1A receptor was observed in the intrarenal vasculature, glomeruli, proximal and distal tubules, and collecting ducts. The AT2 receptor was localized mainly to the glomeruli. The AT1A but not AT2 receptor protein expression was significantly reduced in rats with 10-day systemic Ang II–induced hypertension. In both 7-day 2K1C and 3-day 2K1W rats, the AT1A receptor was significantly reduced in ischemic and contralateral kidneys compared with sham-operated control rats. Reduction in AT2 receptor expression was observed only in the ischemic kidneys in 2K1C and 2K1W renal hypertensive rats. These results demonstrate that the AT1A receptor is widely distributed in the glomerulus and all other nephron segments of the rat kidney. Renal AT1A but not AT2 receptor protein is downregulated in rats with Ang II–induced hypertension. In renal hypertensive rats, the AT1A receptor is bilaterally downregulated and the AT2 receptor is downregulated only in the ischemic kidney. (Hypertension. 1999;33:96-101.)

Key Words: receptors, angiotensin II ▪ immunohistochemistry ▪ kidney ▪ hypertension, renal

The kidney is an important target organ for angiotensin II (Ang II), which plays a critical role in the regulation of blood flow, glomerular filtration, tubular sodium reabsorption, and renin secretion. The 2 subtypes of the AT1 receptor (AT1A and AT1B) are encoded by separate genes on different chromosomes. Their cDNAs have 91% identity in their nucleotide sequences within the coding regions but exhibit only 60% homology in the 5' and 3' untranslated regions. Both AT1A and AT1B receptor mRNAs are present in the kidney, with the AT1A receptor as the predominant isoform, and their mRNA expression is differentially regulated. However, the intrarenal localization of the AT1 receptor subtype protein remains unknown largely because of the lack of specific ligand for different AT1 receptor isoforms. The AT2 receptor is expressed in the adult kidney at a much lower level and is localized mainly to the glomerular mesangial cells. The majority of studies suggest that the renal actions of Ang II are mediated primarily by the AT1 receptor, except for the regulation of electrolyte excretion. Activation of renal AT1 receptors has been reported to regulate pressure-natriuresis, to cause vasodilation in the preglomerular afferent arteriole, and to stimulate renal nitric oxide production.

Both 2-kidney, 1-clip (2K1C) (Goldblatt) and 2-kidney, 1-figure-of-8-wrap (2K1W) (Grollman) renal hypertension are renin dependent. Differential regulation of renal AT1 subtype receptor mRNA in renal hypertension has been reported in 2K1C rats. However, regulation of the AT2 receptor expression in renal hypertension is unknown. The present study is designed to investigate the distribution and regulation of renal subtype AT1A and AT2 receptor protein in Ang II–dependent hypertension in rats induced by systemic Ang II infusion and rats in the acute phase of renal hypertension (either 7-day 2K1C or 3-day 2K1W hypertension).

Methods

All animal procedures were conducted with the approval of the Animal Research Committee of the University of Virginia School of Medicine.

Characteristics of Polyclonal Antisera

The peptide (213 AYEIQKNKPRNDD317), a part of the third cytoplasmic domain of the rat AT1A receptor, was cross-linked to thyroglobulin and used to generate a specific anti–AT1 receptor polyclonal antiserum in the rabbit. This antiserum recognized a specific protein with an estimated molecular mass of 60 kd, consistent with the predicted molecular weight of the AT1A receptor. The specificity of the antiserum was further supported by the single band of 60 kd detected from the AT1A receptor–rich adult (12-week old) rat liver but not from the AT1B receptor–rich adult (12-week old) rat pituitary gland (Pel-Freez;
Rogers, Ariz) by immunoblot analysis, performed as described in the
following section (Figure 1, left); a significant decrease in the
enzyme-linked immunosorbent assay response (10 ng of the peptide
used as antigen target) after preadsorption challenge of the antiserum
with the peptide at 120 μg/mL diluted antiserum, performed as
described5 (Figure 1, right). The polyclonal antiserum to the AT₁ receptor
was raised in the rabbit against a synthetic peptide sequence
(MKDNFSFAATSRNITSS) derived from the rat AT₁ receptor,
and its specificity has been documented.4,16,17

Animal Preparations
Female Sprague-Dawley rats (200 to 220 g body weight, Hilltop
Laboratory Animals; Scottsdale, Pa) were randomly allocated to the
following groups: 2K1C hypertension and a sham-operated control
group; 2K1W hypertension and a sham-operated control group; and
Ang II–induced hypertension and a vehicle-infused control group.3
After rats were anesthetized with pentobarbital sodium (50 mg/kg
body wt intraperitoneally), the left kidneys were exposed by a
midline laparotomy in the first 2 groups. In the first group, the left
renal artery was clipped by placement of a solid U-shaped silver clip.
The effective clipping of the left renal artery was confirmed by a
70% to 75% reduction in renal cortical blood flow as determined by
a laser Doppler probe (superficial probe, type C, Advance Co, Ltd)
coupled to a laser Doppler flowmeter (ALF 21D Dual Channel
Flowmeter, Advance Co, Ltd). In the second group, the left kidney
was tautly wrapped in a figure-of-8 pattern around the kidney to
compress the poles and body with 2.0 silk thread (Ethicon Inc). All
rats in the sham control groups received identical surgical treatment

Figure 1. Characteristics of polyclonal antiserum against rat
AT₁A subtype receptor. Left, Representative immunoblot of AT₁A
receptor protein from liver (lane 1) and pituitary gland (lane 2) of
adult (12-week-old) rat (40 μg of total protein per lane). Right,
Summary of the enzyme-linked immunosorbent assay response
after preadsorption challenge of AT₁A receptor antiserum with
peptide immunogen. n=3, **P<0.01 vs antiserum.

Figure 2. Light photomicrographs of frozen sections of rat kidneys immunostained for
AT₁A receptor expression. In normal rat kidney, positive immunoreactivity was observed
in glomeruli, proximal and distal tubules, and collecting ducts (A, B). Macula densa
cells were strongly stained (A, arrow). Positive immunostaining was also detected in
intrarenal vasculature (A), including afferent arteriole (C, arrow) and interlobar artery (D).
No immunoreactive signal was observed in consecutive sections incubated with preab-
sorbed antiserum (E). Systemic Ang II infusion apparently decreased the intensity of
immunostaining (F). AT₁A receptor immunoreactivity also was notably decreased in
clipped (G) and unclipped (H) kidneys of 2K1C and in wrapped (I) and unwrapped (J)
apart from clip placement or renal wrap. In the third group, Ang II (25 ng/min in lactated Ringer’s solution) or vehicle was infused into the jugular vein through mini-osmotic pumps (model 2002, Alza). After 3 days (for 2K1W and sham-operated control rats), 7 days (for 2K1C and sham-operated control rats), or 10 days (for Ang II–infused or vehicle-infused rats), systolic arterial blood pressure was measured by a tail-cuff photosensor (Rat Tail Monometer-Tachometer system, Natsume model KN-210, Peninsula Laboratories). The kidneys were quickly perfused and then harvested for immunohistochemical studies (n = 3/group) and Western blot analysis (n = 7/group) from rats deeply anesthetized with pentobarbital sodium (80 mg/kg body wt intraperitoneally).

Light Microscopic Immunohistochemistry
Indirect immunoperoxidase immunohistochemistry was performed as previously described.1,4,16 Frozen sections (10 to 12 μm) were incubated with one of the following, diluted 1:500 in 1.5% normal goat serum and 0.5% nonfat dry milk in phosphate-buffered saline: AT1A (7.7 protein mg/mL) or AT2 (3.1 mg protein/mL) receptor primary antiserum; AT1A or AT2 receptor primary antiserum preabsorbed against its respective synthetic peptide antigen. For preabsorption, antiserum was incubated overnight at 4°C with its respective peptide immunogen at 120 μg/mL (for AT1A) or 50 μg/mL (for AT2) diluted antiserum. Immunoreactive signal was detected with an avidin-biotin immunoperoxidase reaction (Vectastain ABC kit, Vector Laboratory) and visualized by exposure to diaminobenzidine for 40 seconds.

Western Blot Analysis
Western blot analysis was performed as previously described.1,4,16 The nitrocellulose membrane was incubated with the AT1A or AT2 receptor antibody (both 1:1000 dilution in Tris-buffered saline with 5% nonfat dry milk and 0.1% Tween 20) in random order. Blots were then incubated with peroxidase-conjugated donkey anti-rabbit secondary antibody (1:5000 dilution, Amersham, Arlington Heights, Ill.). The immunoreactivity was visualized with an enhanced chemiluminescence Western blotting detection kit (Amersham). Membranes were stripped between incubations with different antibodies in a Tris-buffered solution containing 2% sodium dodecyl sulfate and 100 mmol/L β-mercaptoethanol at 50°C.

Quantitative assessment of band densities was performed by scanning densitometry (ImageQuant, Molecular Dynamics). Statistical analysis was performed with a Macintosh StatView program (Abacus Concepts). All data were expressed as mean ± SEM. Comparisons were made with either the unpaired t test or 1-way ANOVA followed by Fisher protected least significant difference test for multiple comparisons when appropriate. A value of P < 0.05 was considered statistically significant.

Results
The 2K1C and 2K1W animals exhibited significantly increased systolic blood pressure compared with their respective sham-operated controls (166 ± 6 vs 95 ± 4 mm Hg in 2K1C, P < 0.01; 148 ± 2 vs 104 ± 3 mm Hg in 2K1W, P < 0.01). Ang II–infused rats also had elevated blood pressure compared with vehicle-infused controls (159 ± 9 vs 100 ± 7 mm Hg, P < 0.01).

Intrarenal Distribution of AT1A and AT2 Subtype Receptors
In the frozen sections of normal rat kidney, intense immunoreactivity for the AT1A receptor was observed in glomeruli, proximal and distal tubules, and collecting ducts (Figure 2, A and B). The macula densa cells were strongly stained (Figure 2, A). Positive immunostaining also was observed in the intrarenal vasculature (Figure 2, A), including juxtaglomerular cells (Figure 2, C) and interlobular artery (Figure 2, D).
No specific staining was observed in consecutive sections incubated with preabsorbed antiserum (Figure 2, E). Notably, less staining than normal control rats was observed in the kidney of the Ang II–infused rat (Figure 2, F) and both ischemic and contralateral kidneys of the 2K1C (Figure 2, G and H) and 2K1W (Figure 2, I and J) rats, respectively.

The immunoreactive signal for the AT2 receptor was present mainly in the glomeruli and also lightly in the proximal tubules of the normal rat kidney (Figure 3, A). No specific staining was observed in consecutive sections incubated with preabsorbed antiserum (Figure 3, B). The AT2 receptor staining was absent in the clipped kidney of the 2K1C rat (Figure 3, C) and apparently decreased in both the glomeruli and tubules in the wrapped kidney of the 2K1W rat (Figure 3, D). However, both the glomeruli and tubular epithelium were positively stained in the contralateral (nonischemic) kidneys of the 2K1C (Figure 3, E) and 2K1W rats (Figure 3, F). No apparent change from the normal rat in the renal AT2 receptor staining was observed in the Ang II–infused rat (Figure 3, G).

Western Blot Analysis of Renal AT1A and AT2 Subtype Receptors
In both 2K1C and 2K1W renal hypertensive rats, the AT1A receptor protein was significantly reduced in the ischemic and contralateral kidneys compared with sham-operated control rats (n=7/group, P<0.01) and 2K1W (n=7/group, P<0.05) rats (Figure 4). No significant change from the normal rat in the renal AT2 receptor protein was observed between Ang II–infused and vehicle-infused rats (n=7/group, P>0.05, Figure 5).

Discussion
Because all known AT1 receptor ligands bind with equivalent affinities to AT1A and AT1B receptor subtypes, the distribution of the AT1 subtype receptor in the kidney has been difficult to evaluate with ligand binding and autoradiographic techniques. Moreover, previous efforts to localize the renal AT1 receptor protein failed to distinguish AT1A from AT1B receptor subtypes because the anti-peptide antibodies used were raised against shared amino acid sequences between AT1A and AT1B receptor subtypes.18 In this study, AT1A subtype receptor protein was localized to the glomeruli and all other nephron segments of rat kidney by immunohistochemistry with the use of a specific anti–AT1A receptor antibody. Our results are in good agreement with the expression pattern of renal AT1A receptor mRNA in that the AT1A receptor subtype is the major (>80%) isoform of AT1 receptor mRNA detected in all nephron segments except the glomerulus, where AT1A and AT1B mRNA are expressed in similar proportions.7 Renal AT1A receptor expression in the afferent and efferent arterio-

**Figure 4.** Representative immunoblot (left) and summary of densitometry quantitation (right) of AT1A receptor protein in 3 groups of hypertensive rats (40 μg of total kidney protein per lane, n=7/group). Top, AT1A receptor from both contralateral (lanes 4, 5, and 6, CTL) and clipped (lanes 7, 8, and 9, CLP) kidneys of 2K1C rats was significantly less than that of sham-operated control rats (lanes 1, 2, and 3, C). Middle, AT1A receptor from both contralateral (lanes 4, 5, and 6, CTL) and wrapped (lanes 7, 8, and 9, WRP) kidneys of 2K1W rats is significantly less than sham-operated control rats (lanes 1, 2, and 3, C). Bottom, Renal AT1A receptor from Ang II–infused hypertensive rats (lanes 4, 5, and 6, Ang II) was significantly decreased when compared with vehicle-infused control rats (lanes 1, 2, and 3, C). *P<0.05, **P<0.01 vs C. Migration and size of molecular weight markers (MW, values ×103) are at left. Approximate molecular mass of AT1A receptor is 60 kd.
lar smooth muscle and in the glomerular mesangial cells was recently demonstrated indirectly by immunohistochemical localization of β-galactosidase in the heterozygous AT1A receptor of mutant mice, generated by a targeted replacement of the AT1A receptor loci by the lacZ gene. However, whether there is any tubular localization of the AT1A receptor was not reported. Ang II influences single-nephron glomerular filtration rate by modifying glomerular vascular tone, inducing contraction of mesangial cells, and reducing glomerular ultrafiltration coefficient. Ang II regulates sodium and bicarbonate transport in proximal tubules; modulates sodium, chloride, and bicarbonate transport in distal tubules; and modifies H+/ATPase activity and amiloride-sensitive sodium transport in the cortical collecting ducts. Our findings provide structural evidence for the possible role of the AT1A receptor in these Ang II–mediated renal effects.

Renal hypertension produced by unilateral renal artery clipping or renal compression is associated with increased renin release and Ang II production from the ischemic kidney (also unpublished data). The current findings provide direct evidence for the bilateral downregulation of the renal AT1A subtype receptor protein in rats with acute-phase 2K1C and 2K1W renal hypertension. Renal AT1A receptor mRNA expression is decreased after systemic administration of a nonpressor dose of Ang II and in the clipped kidney 2 days after unilateral renal artery clipping. These results are in agreement with the notion that increased Ang II exerts a negative regulatory effect on the renal expression of the AT1A receptor.

This study demonstrated differential regulation of the renal AT1 receptor in renal hypertension. The AT1 receptor was downregulated only in the ischemic kidneys of 2K1C and 2K1W hypertensive rats, whereas renal AT1 receptor expression was not significantly affected in Ang II–induced hypertensive rats. These results suggest that increased Ang II itself may not be directly responsible for this downregulation. Hypoperfusion and/or other local changes secondary to reduced blood supply to the clipped/wrapped kidney are possible causes of suppressed AT2 receptor expression in the ischemic kidneys of renal hypertensive rats. A lack of inducible AT2 receptor expression in cultured rat glomerular mesangial cells from stroke-prone spontaneously hypertensive rats and aortic vascular smooth muscle cells from spontaneously hypertensive rats has been described. The AT2 receptor not only causes vasodilation in the preglomerular afferent arteriole but also exerts an antiproliferative effect and induces apoptosis. Recently, accelerated renal interstitial fibrosis and collagen deposition has been observed in adult AT2 receptor null mutant mice during unilateral ureteral obstruction. Thus, decreased AT2 receptor expression in the ischemic kidney may aggravate renal ischemia and promote cell proliferation/fibrosis, possibly augmenting kidney damage in renal hypertension.
In summary, we localized the AT1A subtype receptor protein to glomeruli, proximal and distal tubules (including macula densa), and collecting ducts, as well as the intrarenal vasculature. Renal AT1A but not AT2 receptor protein was downregulated in Ang II–induced hypertension. Expression of AT1A and AT2 receptor protein in the kidney was differentially regulated in renal vascular hypertensive rats: The AT1A receptor was bilaterally downregulated, whereas the AT2 receptor was downregulated only in the ischemic kidney. Differential regulation of angiotensin receptor subtypes may play a role in the pathophysiology of renal hypertension.

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