Intrarenal dopamine modulates progressive angiotensin II-mediated renal injury

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Yang S, Yao B, Zhou Y, Yin H, Zhang MZ, Harris RC. Intrarenal dopamine modulates progressive angiotensin II-mediated renal injury. Am J Physiol Renal Physiol 302: F742–F749, 2012. First published December 14, 2011; doi:10.1152/ajprenal.00583.2011.—It is well-recognized that excessive angiotensin II (ANG II) can mediate progressive renal injury. Previous studies by us and others have indicated that dopamine may modulate actions of ANG II in the kidney. The current studies investigated whether altering intrarenal dopamine levels affected ANG II-mediated renal fibrosis. We utilized a model of increased intrarenal dopamine, catechol-O-methyl-transferase knockout (COMT KO) mice, which have increased kidney dopamine levels due to deletion of a major intrarenal dopamine-metabolizing enzyme. In wild-type mice, chronic ANG II infusion increased renal expression of both of the major dopamine-metabolizing enzymes, COMT and monoamine oxidase. After 8 wk of ANG II infusion, there were no significant differences in blood pressure between wild-type and COMT KO mice. Compared with wild-type, COMT KO mice had decreased albuminuria and tubulointerstitial injury. In response to ANG II infusion, there was decreased expression of both glomerular and tubulointerstitial injury markers (fibronectin, connective tissue growth factor, fibroblast-specific protein-1, collagen I, podocyte vascular endothelial growth factor) in COMT KO mice. We recently reported that ANG II-mediated tubulointerstitial fibrosis is mediated by src-dependent epidermal growth factor receptor (EGFR) activation. In aromatic l-amino acid decarboxylase knockout (AADC KO) mice, a model of intrarenal dopamine deficiency due to selective proximal tubule AADC deletion, which inhibits intrarenal dopamine synthesis, ANG II infusion further increased expression of p-src and pTyrs845-EGFR. In contrast, their expression was markedly attenuated in COMT KO mice. These results demonstrate a role for intrarenal dopamine to buffer the detrimental effects of ANG II upon the kidney.

Dopamine’s cellular actions are mediated by signaling through G protein-coupled seven transmembrane receptors. In mammals, there are five known renal dopamine receptors, which are divided into two subclasses: D1-like and D2-like receptors. D1-like receptors (D1 and D5) are coupled to Gs and stimulate adenylyl cyclase. D2-like receptors (D2, D3, and D4) are coupled predominantly to Gi.

In the mammalian kidney, dopamine receptor activation inhibits both proximal and distal solute and water transport (1, 4). In addition to its role in regulation of salt and water excretion and blood pressure, dopamine may be protective to the kidney under pathological conditions. Both D1-like receptors and D2-like receptors have been reported to protect against acute kidney injury (23, 32). A recent randomized controlled trial indicates that donor pretreatment with dopamine limited cold-ischemia-induced kidney injury following transplantation (25). One mechanism underlying dopamine kidney protection may be its antioxidant effect (35).

Angiotensin II (ANG II), mediated through AT1 receptors, stimulates reabsorption in both proximal and distal nephrons. Thus, dopamine and ANG II appear to serve counterregulatory functions in the kidney (5, 15, 16). In this regard, dopamine inhibits renal renin expression (38) and inhibits ANG II-mediated tubule function and AT1 expression (9, 21, 28, 40). The goal of the present studies was to determine the potential role of intrarenal dopamine to modulate the effects of ANG II excess on renal function and development of progressive injury.

METHODS

Animals. Animal experiments were performed in accordance with the guidelines of the Institutional Animal Care and Use Committee (IACUC) of Vanderbilt University. Animal protocol was reviewed and approved by IACUC of Vanderbilt University. All mice used in the studies were on a 129J/sv background. Wild-type and COMT knockout (KO) mice were obtained from Dr. Maria Karayiorgou at Rockefeller University (17). AADC floxed mice were generated in our laboratories, backcrossed onto the 129J/sv background for 10 generations, and then crossed with 129J/sv γ-GT Cre mice (40). All mice were genotyped before use. ANG II infusion induced more severe kidney injury in unilaterally nephrectomized (UNX) animals than in animals with two kidneys (22). Therefore, in our studies, the UNX mice were treated with ANG II (BACHEM) at a dose of 1.4 mg·kg−1·day−1 (COMT KO and corresponding wild-type mice) through subcutaneous osmotic minipumps (model 2004, Alzet) (20) and killed after 8 wk. Our preliminary experiment found that most of UNX AADC KO mice treated with 1.4 mg·kg−1·day−1 ANG II died within 4 wk after initiation of ANG II infusion. Therefore, a reduced ANG II dose (0.9 mg·kg−1·day−1) and shortened experiment period (4 wk) were employed for AADC KO and corresponding wild-type mice.

The renal selective dopamine precursor gludopa was synthesized in the Chemical Synthesis Core, Vanderbilt Institute of
Chemical Biology. Gludopa at a dose of 5 or 10 mg·kg⁻¹·day⁻¹ was administered through subcutaneous osmotic minipumps (model 2001). Urine samples were collected by using the Créde technique and stored at −80°C until use. Urinary ANG II was measured by GC/electron capture/negative chemical ionization mass spectrometry assay as previously described (40).

Measurements of blood pressure and albuminuria. Systolic blood pressure (SBP) was measured in conscious, trained mice at room temperature using a tail-cuff monitor (BP-2000 BP Analysis System, Visitech System). Twenty-four-hour urine was collected from individually caged mice using polycarbonate metabolic cages. Urinary albumin and creatinine excretion were determined using Alburell-M kits (Exocell, Philadelphia, PA).

Measurement of kidney dopamine. Dopamine was measured by HPLC coupled with electrochemical detection by the Neurochemistry Core Laboratory at Vanderbilt University’s Center for Molecular Neuroscience Research.

Antibodies. Rabbit antibodies against AADC (AB136) and collagen I were purchased from Millipore. Goat anti-COMT was from Novus. Rabbit anti-fibroblast-specific protein-1 (FSP-1) was a gift from Dr. Eric Neilson (Vanderbilt University), rabbit anti-human fibronectin was from Sigma, rabbit anti-Mas was from Alomone (AAR-013), monoclonal anti-angiotensinogen (AGT) was from ABBOTT (catalog no. 200551), and rabbit anti-p-Src was from Cell Signaling Technology. Rabbit anti-p-epidermal growth factor receptor (EGFR) (Tyr848) and EGFR, MAO A/B, goat anti-connexin tissue growth factor (CTGF), and NADPH oxidase 1 (NOX-1) were from Santa Cruz Biotechnology.

Immunohistochemistry and Western analysis and quantitative image analysis. Under deep anesthesia with Nembutal (70 mg/kg ip, Abbot Laboratories), mice were exsanguinated with aortic cannula. One kidney was removed for Western blot analysis. The other kidney was perfused as previously reported (37). The fixed kidneys were dehydrated through a graded series of ethanols, embedded in paraffin, sectioned at 4-μm thickness, and mounted on glass slides. The slides were deparaffinized, rehydrated, and stained with different antibodies, as previously described (18). Based on the distinctive density and color of immunostaining in video images, the number, size, and position of stained cells were quantified using the BIOQUANT true-color windows system (R & M Biometrics, Nashville, TN) as previously described (36, 37).

Whole blot analysis was carried out as described previously (7, 26).

Table 1. Primers for dopamine and angiotensin systems in the cultured proximal tubule epithelial cells

<table>
<thead>
<tr>
<th>Primers</th>
<th>Forward</th>
<th>Reverse</th>
<th>Product</th>
</tr>
</thead>
<tbody>
<tr>
<td>AADC</td>
<td>GTTTGCGTTCAGCGGTTCTG</td>
<td>CGAAGACCGAGTGTAGTTA</td>
<td>200 bp</td>
</tr>
<tr>
<td>COMT</td>
<td>CGCAAGCTTTCATCAGCT</td>
<td>TCTGACTTCTCTTCTCACG</td>
<td>143 bp</td>
</tr>
<tr>
<td>MAO-A</td>
<td>GCAAACAGCTGAGTTGCTA</td>
<td>GCCACAGAATGAAAAACCG</td>
<td>224 bp</td>
</tr>
<tr>
<td>MAO-B</td>
<td>GTGTTGCTGGAGCGAGCTG</td>
<td>TCTAGTCCGCTGAAAGAACACT</td>
<td>723 bp</td>
</tr>
<tr>
<td>DAT</td>
<td>ATCATGCGGACACTTTCTAC</td>
<td>CATATGGGCAAGAAGAACGGTG</td>
<td>95 bp</td>
</tr>
<tr>
<td>D1</td>
<td>ATCATGCGGACACTTTCTAC</td>
<td>GCCCGCTGAGAGCCACCCAG</td>
<td>230 bp</td>
</tr>
<tr>
<td>D2</td>
<td>ATCATGCGGACACTTTCTAC</td>
<td>AGCCGCTGAGAGCCACCCAG</td>
<td>208 bp</td>
</tr>
<tr>
<td>D3</td>
<td>ATCATGCGGACACTTTCTAC</td>
<td>GATTTGCGTGGCGGTTGTA</td>
<td>310 bp</td>
</tr>
<tr>
<td>D4</td>
<td>ATCATGCGGACACTTTCTAC</td>
<td>TCCGGACCCGAGCTCTGAA</td>
<td>202 bp</td>
</tr>
<tr>
<td>D5</td>
<td>ATCATGCGGACACTTTCTAC</td>
<td>ATCTGTCGCTGAGATT</td>
<td>221 bp</td>
</tr>
<tr>
<td>AT1a</td>
<td>GACATGCTGTTCGGTCTGTC</td>
<td>GGAAGAGCTTTGCTATG</td>
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</tr>
<tr>
<td>AT1b</td>
<td>GGAGGATCTGCTGCTGATCGT</td>
<td>GGAAGAGCTTTGCTATG</td>
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</tr>
<tr>
<td>AT2</td>
<td>ATGCTGCTGCTGCTGATCGT</td>
<td>AGCTGGAAGAAAGAAGGAGA</td>
<td>454 bp</td>
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<tr>
<td>Mas</td>
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<td>GAGTGGAGGAAAGAAG gag</td>
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<tr>
<td>AGT</td>
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<td>CGAAGATCTGCTGCTGATCGT</td>
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</tr>
<tr>
<td>ACE</td>
<td>ATGCTGCTGCTGCTGATCGT</td>
<td>CGAAGATCTGCTGCTGATCGT</td>
<td>340 bp</td>
</tr>
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<td>ACE2</td>
<td>ATGCTGCTGCTGCTGATCGT</td>
<td>CGAAGATCTGCTGCTGATCGT</td>
<td>431 bp</td>
</tr>
</tbody>
</table>

AADC, aromatic l-amino acid decarboxylase; COMT, catechol-O-methyltransferase; MAO, monoamine oxidase; DAT, dopamine transporter; AT1a, and AT1b, angiotensin II receptors type 1a and 1b; AT2, angiotensin II receptor type 2; Mas, Ang I-7 receptor; AGT, angiotensinogen; ACE, angiotensin-converting enzyme.
note, urinary ANG II excretion was markedly decreased in COMT KO mice, further indicating intrarenal effects of altering renal dopamine metabolism.

Intravenous infusion of ANG II in humans acutely reduces urinary dopamine levels (13), but the role of ANG II to modulate expression of components of intrarenal dopamine production has not been systematically examined. As indicated in Fig. 3A, chronic ANG II infusion did not alter renal expression of AADC, the enzyme that converts L-DOPA to active dopamine, in either wild-type or COMT KO mice. In contrast, it increased expression of COMT and MAO, the two major intrarenal dopamine-metabolizing enzymes in wild-type mice. In COMT KO mice, chronic ANG II infusion also increased expression of MAO. ANG II decreased renal dopamine in wild-type mice, but in COMT KO mice, intrarenal dopamine levels were not statistically different than nontreated mice. These results suggest that COMT, but not MAO, plays a major role in ANG II-mediated reduction of kidney dopamine production.

After 8 wk of ANG II infusion (1.4 mg·kg⁻¹·day⁻¹), there were no significant differences in blood pressure between wild-type and COMT KO mice (SBP: 185 ± 3 vs. 186 ± 2 mmHg of ANG II-infused wild-type mice, n = 8). As indicated in Fig. 4B, ANG II infusion led to markedly less proteinuria in UNX COMT KO mice than in UNX wild-type mice. At baseline, albuminuria was similar in wild-type and COMT KO mice (33 ± 3 vs. 35 ± 2 μg/day, n = 6). In wild-type mice, UNX for 8 wk led to increased albuminuria, which was further augmented by ANG II infusion (μg/day: UNX: 256 ± 79, P < 0.01 vs. control; UNX + ANG II: 8,110 ± 1,437, P < 0.01 vs. UNX, n = 6 in each group; Fig. 4). In contrast, COMT KO mice had minimal increases in albuminuria with UNX (45 ± 6) and significantly less albuminuria with ANG II than in wild-type (UNX + ANG II: 2,697 ± 632, P < 0.01, n = 6; Fig. 4).

Histologic analysis indicated that ANG II infusion led to interstitial fibrosis, as indicated by Masson trichrome staining, and glomerular sclerosis as indicated by silver staining in wild-type mice, and both glomerular and tubulointerstitial fibrosis were significantly attenuated in COMT KO mice (Fig. 5). Immunohistochemistry showed that in COMT KO mice, there was attenuation of the ANG II-induced increases...
and phospho-845YEGFR expression were inhibited in COMT in response to chronic ANG II infusion, both src activation and src-dependent phosphorylation of tyrosine residue 845 (6). Therefore, it was noteworthy that in the current studies, src activation of src kinase, with activation of EGFR by phosphorylation of tyrosine 845 in the EGFR cytoplasmic domain, which is known to be a target of src phosphorylation (6). Therefore, it was noteworthy that in the current studies, src activation and 845YEGFR expression were significantly decreased in COMT KO mice and were increased in AADC KO mice. Whether this is secondary to the known effect of dopamine to decrease AT1 receptor expression in the proximal tubule (9, 27, 28) or to direct antioxidant effects of dopamine (2, 29–31) will require further studies.

COMT is a major intrarenal dopamine-metabolizing enzyme, and COMT KO mice have increased intrarenal and urinary dopamine levels due to the absence of COMT metabolism of dopamine. However, plasma dopamine concentrations are similar between wild-type and COMT KO mice (19, 38). Previous studies indicate that ANG II may regulate the intrarenal dopaminergic system at multiple levels. ANG II has been reported to inhibit dopamine uptake and AADC activity but to stimulate MAO activity in kidney cortex through activation of AT1 receptors (10–12). Nitecapone, a peripheral inhibitor of COMT, induces dopamine-dependent natriuresis (14). However, it has been suggested that entacapone, another inhibitor of COMT, protected from

Fig. 2. Urinary ANG II excretion was inhibited by activation of the intrarenal dopaminergic system. Gludopa, a renal specific dopamine precursor, inhibited urinary ANG II excretion in wild-type mice. Compared with wild-type mice, urinary ANG II excretion was reduced in COMT knockout (KO) mice. *P < 0.05 vs. control wild-type mice, n = 4 in each group.

in interstitial fibronectin, collagen I, and the fibroblast marker FSP-1 observed in wild-type mice (Fig. 6A). Podocyte vascular endothelial growth factor expression was increased in response to ANG II in wild-type mice, but this increase was markedly attenuated in COMT−/− mice. Similarly, CTGF expression increased in both glomeruli and tubules in response to ANG II but there were minimal increases in COMT−/− mice. Immunoblotting confirmed these differences in renal expression of fibronectin, CTGF, and FSP-1 (Fig. 6B).

We and others reported previously that the tubulointerstitial injury induced by chronic ANG II infusion is predominantly mediated by EGFR transactivation (6, 20), and we previously found that this effect was mediated by reactive oxygen species activation of src, which activates EGFR by src-dependent phosphorylation of tyrosine residue 845 (6). In response to chronic ANG II infusion, both src activation and phospho-845YEGFR expression were inhibited in COMT KO mice compared with wild-type mice (Fig. 7A). Also of interest, NOX-1 expression increases in response to ANG II infusion were blunted in COMT KO mice (Fig. 7B). We previously described increased renal injury in response to ANG II in mice with selective intrarenal dopamine production due to genetic deletion of AADC (40). In these mice, expression of phospho-src and phospho-845YEGFR was increased in response to chronic ANG II administration compared with that in wild-type mice (Fig. 7C).

Fig. 3. ANG II activated renal dopamine-metabolizing system and decreased renal dopamine levels. A: chronic ANG II infusion led to increases in expression of MAO and COMT. B: at baseline, renal dopamine levels were higher in COMT KO mice than in wild-type. ANG II infusion decreased renal dopamine levels in wild-type but not COMT KO mice (*P < 0.05 vs. control wild-type, n = 4 in each group).
ANG II-induced inflammation and renal injury due to its anti-oxidant effect but not due to a change in renal dopaminergic tone (19). The possible explanation for the lack of effect of entacapone on renal dopaminergic tone is that MAO activity may be predominant in metabolism of dopamine in these transgenic rats. It is well-known that dopamine and ANG II can each antagonize the activity of the other hormone, but this observed increase in protein expression of the major intrarenal pathways of dopamine metabolism in response to ANG II represents a previously undescribed mechanism by which ANG II may decrease dopamine activity in the kidney.

We previously reported that dopamine may indirectly decrease ANG II production by inhibiting renal renin production (38–40). In the current studies, dopamine decreased AGT expression in cultured proximal tubule cells, and gludopa decreased urinary ANG II production. Administration of dopamine to proximal tubule cells also increased mRNA for ACE2, the mediator of production of Ang1–7, which serves as a counterregulatory hormone to AT1-mediated responses. In addition, previous studies demonstrated that dopamine will increase activity of AT2 receptors (24), which can also antagonize ANG II’s actions via AT1 receptors. Therefore, in addition to direct inhibition of angiotensin expression and AT1 expression and activity, dopamine’s antago-

Fig. 4. ANG II-mediated albuminuria was significantly attenuated in COMT KO mice. A: SDS-PAGE gel electrophoresis indicates significantly less urinary protein excretion in ANG II-treated unilaterally nephrectomized (UNX) COMT KO mice than in the ANG II-treated UNX wild-type mice. Loading volume was determined using urinary creatinine as an internal control. B: 24-h urinary albumin excretion (UAE) was significantly lower in ANG II-treated UNX COMT KO mice than in ANG II-treated UNX wild-type mice (*P < 0.01 vs. wild-type control; †P < 0.01 vs. corresponding UNX mice; ‡P < 0.01 vs. ANG II-treated UNX wild-type mice, n = 6 in each group).

Fig. 5. ANG II-mediated renal injury was markedly attenuated in COMT KO mice. ANG II infusion led to significantly less interstitial fibrosis (A) and glomerular sclerosis (B) in COMT KO mice than in wild-type mice (*P < 0.01 vs. wild-type, n = 4).
nism of AT1 may also be mediated in part by upregulating angiotensin-mediated signaling pathways that antagonize AT1-mediated signaling.

In summary, these studies provide evidence for the important counterregulatory role that the intrarenal dopaminergic system plays to counteract the effects of ANG II signaling through AT1 receptors (Fig. 8). In addition to previous studies indicating that dopamine can decrease renin production and decrease AT1 receptor expression in the kidney, the current studies indicate that dopamine decreases intrarenal angiotensin production. Furthermore, in the face of chronic ANG II excess, increased intrarenal dopamine prevents increases in expression of profibrotic signaling pathways and it decreases glomerular and tubulointerstitial injury.

Fig. 6. ANG II-mediated elevations of mediators and markers of fibrosis were attenuated in COMT KO mice. A: immunohistochemistry showed that ANG II-induced elevations of fibronectin, connective tissue growth factor (CTGF), fibroblast-specific protein-1 (FSP-1), collagen I, and vascular endothelial growth factor (VEGF) were significantly attenuated in COMT KO mice. B: immunoblotting indicated that ANG II-induced increases in fibronectin, FSP-1, and CTGF were attenuated in COMT KO mice.

Fig. 7. ANGII-mediated activation of Src-epidermal growth factor receptor (EGFR) signaling was attenuated by intrarenal dopamine. A: activation of Src and EGFR induced by ANG II was attenuated in COMT KO mice. B: ANG II-induced increases in NADPH oxidase 1 (NOX-1) expression were inhibited in COMT KO mice. Control and ANG II-infused samples were from the same gel. C: activation of Src and EGFR induced by ANG II was augmented in AADC KO mice.
Fig. 8. Intrarenal dopamine antagonizes ANG II-induced renal injury at multiple levels. ANG II induces renal injury through activation of AT1 receptors. Intrarenal dopamine may antagonize ANG II-induced renal injury through inhibition of AT1 expression and decreases in intrarenal ANG II production due to inhibition of expression of both AGT and renin.

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DISCLOSURES
No conflicts of interest, financial or otherwise, are declared by the author(s).

AUTHOR CONTRIBUTIONS

REFERENCES


