Papillary Collecting Tubule Synthesis of Prostaglandin E2 in Dahl Rats

GERALD M. REID, RICHARD G. APPEL, AND MICHAEL J. DUNN

SUMMARY Isolated kidneys of Dahl salt-sensitive rats (DS) excrete sodium less readily than those of Dahl salt-resistant rats (DR). The collecting tubule is an important source of papillary prostaglandin E2 and is a site of significant sodium reabsorption. We cultured renal papillary collecting tubule cells from 5-week-old, prehypertensive DS and DR on a low salt diet and also after 14 weeks of high salt feeding, and we measured prostaglandin E2 synthetic capacity. Unstimulated renal papillary collecting tubule cells from 5-week-old DS produced 62 ± 5% less prostaglandin E2 than did comparable cells from DR (p<0.001). The cells from DS also synthesized less prostaglandin E2 after stimulation with the calcium ionophore A23187 (67 ± 6% of control; p<0.001) or the addition of exogenous arachidonate (74 ± 7% of control; p<0.01). Urinary prostaglandin E2 excretion was also diminished in the 5-week-old DS compared with their salt-resistant counterparts (18.1 ± 1.3 vs 23.9 ± 1.7 ng/24 hr; p<0.025). After high salt feeding, the DS became hypertensive but the DR remained normotensive. Renal papillary collecting tubule cells cultured from these DS continued to produce less prostaglandin E2 than those from control rats, both in the basal state (60 ± 12% of control; p<0.09) and after stimulation with ionophore (62 ± 2% of control; p<0.002). In these older animals, the DS continued to underexcrete prostaglandin E2 compared with the DR (29.7 ± 3.2 vs 42.2 ± 6.1 ng/24 hr; p<0.08). The underproduction of prostaglandin E2 in the papillary collecting tubule of DS may play a role in their inadequate renal natriuretic capacity and contribute to the onset and maintenance of salt-induced hypertension in this strain. (Hypertension 11: 179-184, 1988)

KEY WORDS • Dahl rats • prostaglandin E2 • collecting tubule • cell culture

Dahl salt-sensitive rats (DS) become hypertensive when fed a diet rich in NaCl, whereas Dahl salt-resistant rats (DR) remain normotensive on a high salt diet. The kidney is of major importance in determining this response. Renal cross-transplantation1,2 experiments between DS and DR have shown that salt-sensitivity transfers with the genotype of the donor. Using isolated perfused kidneys, Tobian et al.3 and Maude and Ko4 found that, at similar perfusion pressures, DR kidneys display a greater natriuresis than DS kidneys and that, to quantitatively equalize natriuresis, DS kidneys need a higher perfusion pressure. This defect in natriuresis probably leads to hypertension through a more complex mechanism than simple volume expansion.5 Natriuretic sub-

stances may be released that have diffuse effects on ion transport, leading to increased vascular reactivity.

A number of observations suggest that the synthetic capacity for renal prostaglandin E2 (PGE2) may be different between the DS and DR strains, especially in the medulla. Urinary PGE2 excretion, which is predominantly medullary in origin,6 was decreased in the DS as compared with the DR, at prehypertensive and hypertensive phases.7 The PGE2 content of snap-frozen renal papillae was also less in the DS than in the DR, both before and after high salt diets.8 Renal medullary microsomes from DS also underproduced PGE2 as compared with microsomes from DR.9

Reduction of medullary PGE2 may be important in the defective natriuresis observed in the DS strain. Prostaglandins have a role in promoting natriuresis in the inner medullary collecting duct in volume-expanded rats,10 a situation somewhat analogous to high salt feeding. Indeed, the inner medullary collecting duct may reabsorb up to 6% of the filtered sodium load11 in volume expansion. This segment is a rich source of medullary PGE2,12,13 Microdissected rabbit nephron segments from the medullary collecting tubule synthesize greater amounts of PGE2 than any other segment of the tubule.14
As cells from the renal papillary collecting tubule (RPCT) can be cultured and produce notable amounts of PGE₂, we hypothesized that RPCT cells from DS might express a genetic defect in PGE₂ synthesis as compared with RPCT cells from DR. We tested this hypothesis using cultured RPCT cells from prehypertensive and hypertensive DS and age-matched DR controls. We found significantly lower PGE₂ synthesis by RPCT cells from DS, whether prehypertensive or hypertensive.

Materials and Methods

Animals

Weanling 3½-week-old male DS and DR were purchased from Brookhaven National Laboratories (Upton, NY, USA). They were maintained on 0.3% NaCl rat chow until 5 weeks of age. Initial urine collections and blood pressure measurements were performed, and the animals were either killed for cell culture experiments or begun on 4% NaCl rat chow for a total of 14 weeks. Color-coded 0.3% NaCl and 4% NaCl rat chow were obtained from ICN Biochemicals (Cleveland, OH, USA).

Blood Pressures

Blood pressures for the 5-week-old rats were obtained directly through an intra-aortic catheter with the rats under pentobarbital anesthesia (40 mg/kg i.p.). Indirect tail-cuff measurements obtained with the rats under light ether anesthesia were used after 9 and 14 weeks of 4% NaCl diet. A minimum of four such readings were made for each rat.

Urine Studies

Twenty-four-hour urine samples were collected in metabolic cages at 5 weeks of age and again after 14 weeks of high salt feeding. Sodium azide, 25 mg, and meclofenamate, 250 mg, were added to each urine container, and 2.5% dextrose was added to the drinking water of the young animals to ensure adequate urine volume. Aliquots (1–2 ml) from the urine samples were acidified to pH 3.0 to 3.5 with citric acid, and PGE₂ (2000 cpm, specific activity, 160 Ci/mmol, Amersham Arlington Heights, IL, USA) was added to each sample to determine recovery. The samples were then extracted twice with ethyl acetate, dried, and dissolved in toluene/ethyl acetate/methanol, 60:40:2. Silicic acid (Mallinckrodt, McGaw Park, IL, USA) columns were prepared with toluene/ethyl acetate/methanol, 60:40:20. After an elution with toluene/ethyl acetate, 60:40, PGE₂ was eluted with the 60:40:20 solvent. Recoveries were 50 to 75%. PGE₂ was measured by radioimmunoassay using PGE₂ antisera from Institut Pasteur (Paris, France).

Renal Papillary Collecting Tubule Cultures

RPCT cultures were done on 5-week-old rats or after 14 weeks of high salt feeding. Each culture was obtained from the pooled kidneys of two or three rats from the same strain. The cultures were carefully paired prospectively such that kidney isolations and preparation were done concurrently from groups of DR and DS, all isolation vehicles and media were identical for each parallel culture, and each culture dish contained wells from both DR and DS cultures.

Cell cultures were performed as previously described. Animals were killed, and their kidneys were placed in ice-cold phosphate-buffered saline. Papillae were carefully dissected and minced. After a 2-hour collagenase (CLS II Lot 4176, Worthington, Freehold, NJ, USA) digestion at 37°C, the incubates were exposed to a brief hypotonic shock followed by a 10% albumin density centrifugation. Cells were initially seeded in a Hams F-12–Dulbecco’s minimal essential medium containing 10% fetal bovine serum to facilitate attachment. After 24 hours this medium was changed to a freshly prepared, defined, K-1 epithelial growth medium. After 72 hours of culture, the cells were nearly confluent and were used for biochemical study. The rate of cell growth was similar for the two strains. Culture wells were washed twice and then incubated for 30 minutes in a 5% CO₂, 95% O₂ atmosphere at 37°C in either minimal essential medium alone (basal) or in medium containing a final concentration of 2 µM Ca⁺² ionophore A23187 (Calbiochem-Behring, San Diego, CA, USA) or arachidonic acid (Sigma Chemical, St. Louis, MO, USA), 5 µg/ml. At the end of the incubation, the supernates were aspirated and quickly frozen. The cell cultures were digested in 1 N NaOH and then assayed by the technique of Lowry et al. for total cell protein per well. PGE₂ was measured on unextracted samples by radioimmunoassay.

Medullary Slices

Medullary slices were prepared from DS and DR after 14 weeks of 4% NaCl feeding as described by Lee and Peter. Rats were anesthetized (pentobarbital, 50 mg/kg i.p.), and kidneys were rapidly excised and placed in ice cold Hanks balanced salt solution. Medullae were dissected, and 0.5-mm slices were prepared with a Stadie Riggs tissue slicer (Thomas Scientific, Swedesboro, NJ, USA). Slices were incubated for 20 minutes at 37°C in a Dubnoff shaker (Research Specialties, Richmond, CA, USA) in Krebs-Henseleit bicarbonate-buffered saline with 30 µM arachidonic acid under a 95% O₂, 5% CO₂ atmosphere. PGE₂ was measured on unextracted incubation media using radioimmunoassay. PGE₂ synthetic rates are expressed per milligram of dry weight, which was determined after a 16-hour desiccation at 80°C.

Statistics

All results were analyzed using independent t tests except those for Tables 2 and 3, where statistical calculations were based on randomized block analysis of variance on log-transformed data.

Results

At 5 weeks of age, while fed only the 0.3% NaCl diet, both DS and DR groups were normotensive. Intra-aortic mean pressure was higher in the DS than in
the DR (97.4 ± 2.9 vs 85.3 ± 1.0 mm Hg; p < 0.001). Blood pressures rose in both groups after high salt feeding, more dramatically in the DS strain. After 14 weeks of a 4% NaCl diet, the DS became frankly hypertensive compared with age-matched DR (160 ± 5.8 mm Hg vs 109.8 ± 1.6 mm Hg; p < 0.001).

In the young, prehypertensive animals, the DS excreted less urinary PGE2 than the DR (Table 1), although total urinary volumes were similar between the groups. After high salt feeding, PGE2 excretion increased in both 19-week-old DS and DR. However, DS continued to excrete less PGE2 than DR, again with no difference between the strains in urinary volume (see Table 1).

RPCT PGE2 production was assessed in the basal condition (i.e., unstimulated RPCT cells) and after stimulation with the Ca2+ ionophore A23187 (2 μM) or arachidonic acid (5 μg/ml). The means of replicate wells for each experiment are shown in Table 2. In the basal, as well as in each of the stimulated conditions, the RPCT cells from DS rats synthesized less PGE2, than the cells from DR (expressed as pg/μg cell protein). The cell protein content was similar between each of the paired, cultured strains.

<table>
<thead>
<tr>
<th>Strain</th>
<th>PGE2 (ng/24 hr)</th>
<th>Urine volume (ml/24 hr)</th>
</tr>
</thead>
<tbody>
<tr>
<td>5-week-old</td>
<td></td>
<td></td>
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<tr>
<td>DR (n=22)</td>
<td>23.9 ± 1.7</td>
<td>44.5 ± 2.0</td>
</tr>
<tr>
<td>DS (n=22)</td>
<td>18.8 ± 1.3*</td>
<td>43.6 ± 2.4</td>
</tr>
<tr>
<td>19-week-old</td>
<td></td>
<td></td>
</tr>
<tr>
<td>DR (n=18)</td>
<td>42.2 ± 6.1†</td>
<td>20.1 ± 2.0</td>
</tr>
<tr>
<td>DS (n=17)</td>
<td>29.7 ± 3.2‡</td>
<td>22.7 ± 2.2</td>
</tr>
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</table>

All values are means ± SEM.
* p < 0.025, compared with values for 5-week-old DR; † p < 0.005, compared with values for same strain at 5 weeks of age; ‡ p < 0.08, compared with values for 19-week-old DR.

Discussion

The kidneys of Dahl rats are important determinants of the blood pressure response to a high salt diet.1,2 The DS have defective sodium chloride excretion as compared with the DR strain.3,4 Kidneys from DS also have decreased medullary PGE2 content.5 PGE2, an

In an effort to negate the week-to-week variability in cultured-cell PGE2 synthesis and stimulability, we decided to assess more carefully each weekly experiment to isolate the effect of strain origin alone. For each culture pair, in each of the three assay conditions, the mean production in the DS wells can be expressed as a percentage of the concurrent DR well production. These data are graphically displayed in Figures 1, 2, and 3. When looked at in this way, the DS cells again produced less PGE2 than their DR counterparts in basal and in both stimulated states.

RPCT cultures were obtained from three additional sets of DS and DR animals after 14 weeks of high salt feeding. PGE2 synthetic capacity was tested under both basal and A23187-stimulated conditions for each culture. The means for the six replicate wells in each experiment are shown in Table 3. DS cultures produced less PGE2 than DR in all cultures under both conditions. This observation was statistically significant when the cells were stimulated, but it failed to reach significance in the basal state because of the small number of studies performed. When the data are expressed as a percentage of PGE2 production for each culture, the DS cells synthesized 60 ± 12% of DR production in the basal state (p < 0.09) and 62 ± 2% of DR production after stimulation (p < 0.002). Finally, medullary slices were stimulated with 30 μM arachidonic acid from five 19-week-old rats of each strain, after the high salt diet. The DS produced 2551 ± 624 pg PGE2/min/mg dry weight, while the DR produced 4389 ± 578 pg PGE2/min/mg dry weight (p < 0.06).

Table 2. Renal Papillary Collecting Tubule PGE2 Production in 5-Week-Old Rats on Low Salt Intake

<table>
<thead>
<tr>
<th>Experiment no.</th>
<th>Basal</th>
<th>A23187 (2 μM)</th>
<th>Arachidonate (5 μg/ml)</th>
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<tr>
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<tr>
<td>1</td>
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Mean ± SEM 2.0 ± 0.3* 3.2 ± 0.4 14.8 ± 2.8* 22.2 ± 3.9 51.1 ± 14.8* 70.8 ± 22.5

Values are means of six to 18 replicate wells.
* F = 39.16, p < 0.001; † F = 22.52, p < 0.002; ‡ F = 15.37, p < 0.01; compared with respective values for DR.
abundant medullary autacoid, has been shown to inhibit sodium chloride transport in the thick ascending limb of Henle of the mouse and the cortical collecting tubule of the rabbit. There is also indirect evidence, in vivo, of prostaglandin-mediated inhibition of sodium reabsorption in the medullary collecting tubule of the rat.

To investigate the onset phase of hypertension in Dahl rats, we conducted our initial studies in young, prehypertensive animals. Urinary PGE₂ excretion, which is predominantly of renal inner medullary origin, was clearly decreased in the DS compared with the DR, confirming the work of others. We next looked at RPCT cells, cultured from the inner medullary collecting tubules of DS and DR, and showed that RPCT cells derived from DS synthesized less PGE₂ than those from DR. The reduced PGE₂ synthesis (26–38% decrement) was detectable in both basal and stimulated (calcium ionophore or arachidonic acid) conditions. We believe that this finding may represent a genetically inherited difference between the strains and may play a role in the defective natriuresis of the DS kidneys. The inner medullary collecting tubule is the final site for salt transport in the nephron and can play a critical role in determining the final electrolyte content of the urine. In the studies of Stein et al., up to 6% of the filtered sodium load was reabsorbed in this segment in young, volume-expanded rats.

We also felt that a defect in PGE₂ production may persist in older animals and thereby be important in the maintenance phase of salt-induced hypertension in the DS. After 14 weeks of high salt feeding, the DS were hypertensive whereas the DR remained normotensive. At this time, urinary PGE₂ excretion was decreased in...
the DS compared with the DR. RPCT cultures from the DS synthesized 40% less PGE_2 than cultures from the DR in the basal and 38% less in Ca^{2+} ionophore–stimulated experiments. In addition, medullary slices from 19-week-old DS released less PGE_2 after the addition of exogenous arachidonic acid than slices from DR, further supporting the cell culture findings.

Work from the laboratory of Tobian et al. underscores the possible importance of the PGE_2 deficit in the DS strain. When these rats were placed on a high linoleic acid diet, renal papillary PGE_2 content rose and the salt-induced rise in blood pressure was significantly blunted. Linoleic acid is chain-elongated and desaturated to the prostaglandin precursor arachidonic acid. Augmented PGE_2 synthesis in peripheral vasculature as well as the kidney may be important in the decreased salt sensitivity in these studies. It should be noted that both linoleic acid, which increased renal prostaglandins, and linolenic acid, which decreased renal prostaglandins, decreased blood pressure in rats with renovascular hypertension.

Although patients with essential hypertension are a heterogeneous group, renal prostaglandin synthesis may be important in some instances. Urinary PGE_2 excretion studies in these patients have yielded conflicting results. However, there appears to be a group of essential hypertensive patients with markedly subnormal urinary PGE_2 excretion, and these patients may also belong to the subgroup that is highly salt-sensitive. Indeed, in one well-studied case, urinary PGE_2 was extremely low and hypertension was salt-sensitive. Furthermore, linoleic acid diets have lowered blood pressure in essential hypertensive humans, although the mechanism of blood pressure reduction is unknown.

Our results, demonstrating decreased PGE_2 synthesis by the RPCT cells from DS in culture, do not unambiguously indicate which enzymatic step in the prostaglandin biosynthetic sequence may be involved. Critical enzymatic sites that regulate the eventual production of PGE_2 include the activation and deacylation of arachidonic acid in membrane phospholipids (acyltransferase, phospholipase), the cyclooxygenation of arachidonic acid to endoperoxides (cyclooxygenase), and the conversion of endoperoxides to PGE_2 (PGE_2 isomerase). The reduced PGE_2 synthesis in the DS strain in the basal state is consistent with an alteration at any of these sites. The reduced stimulating ability of PGE_2 by the Ca^{2+} ionophore A23187 is consistent with defective phospholipase activation or deficient availability of arachidionate but is also consistent with reduced activity of the enzymes beyond the phospholipase step. Abnormalities of these more distal enzymes are further suggested by reduced PGE_2 synthesis in response to arachidonic acid in the DS-derived RPCT cells, especially from the younger, prehypertensive animals. Evaluation of the PGE_2 isomerase would only be possible with the addition of PGH_1 to the cells in culture. Hence, we must conclude that the defect of PGE_2 synthesis may be at multiple enzymatic steps or could be solely attributable to defective cyclooxygenase or PGE_2 isomerase.

In summary, we postulate that defective medullary PGE_2 synthesis in DS may be one factor leading to a decreased capacity to excrete salt, especially in response to an increased dietary salt load. The decreased salt excretion leads to higher blood pressure, probably through a series of complex interactions. The higher perfusion pressure brings the DS back into salt balance, at the expense of increased systemic pressure. This mechanism may be operative in both the genesis and maintenance phases of salt-induced hypertension in this model.

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