

Administration of Atrial Natriuretic Factor Inhibits Sodium-coupled Transport in Proximal Tubules

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Abstract

The newly discovered peptides extracted from cardiac atria, atrial natriuretic factors (ANFs), when administered parenterally cause renal hemodynamic changes and natriuresis. The nephron sites and cellular mechanism accounting for profound increase in Na^+ excretion in response to ANFs are not yet clarified. In the present study we investigated whether synthetic ANF peptide alters the reabsorption of Na^+ and reabsorption of solutes cotransported with Na^+ in the proximal tubules of rats. Synthetic ANF peptide consisting of 26 amino acids, 4 $\mu\text{g/kg}$ body wt/h, or vehicle in controls, was infused to surgically thyroparathyroidectomized anesthetized rats. After determination of the fractional excretion (FE) of electrolytes (Na^+ , K^+ , P_i , Ca^{2+} , Mg^{2+} , HCO_3^-), the kidneys were removed and luminal brush border membrane vesicles (BBMV) were prepared from renal cortex. Solute transport was measured in BBMV by rapid filtration techniques. Infusion of ANF peptide increased FE_{Na} , FE_{P_i} , and $\text{FE}_{\text{HCO}_3^-}$; but FE_{Ca} , FE_{K} , and FE_{Mg} were not changed. The increase in FE_{Na} was significantly correlated, on the one hand, with increase of FE_{P_i} ($r = 0.9$, $n = 7$; $P < 0.01$) and with increase of $\text{FE}_{\text{HCO}_3^-}$ ($r = 0.89$, $n = 7$; $P < 0.01$). On the other hand, FE_{Na} did not correlate with FE_{K} , FE_{Ca} , or with FE_{Mg} . The Na^+ gradient-dependent uptake of P_i by BBMV prepared from renal cortex of rats receiving ANF infusion was significantly ($P < 0.05$) decreased (-25%), whereas the Na^+ gradient-dependent uptake of L-[^3H]proline and of D-[^3H]glucose or the diffusional uptake of $^{22}\text{Na}^+$ were not changed. ANF-elicited change in FE_{P_i} showed a close inverse correlation with decrease of Na^+ -dependent P_i uptake by BBMV isolated from infused rats ($r = 0.99$, $n = 7$; $P < 0.001$). Direct addition of ANF to BBMV in vitro did not change the Na^+ gradient-dependent P_i uptake. In rats infused with ANF, the rate of amiloride-sensitive Na^+ - H^+ exchange across the brush border membrane (BBM) was significantly ($P < 0.05$) decreased (-40%), whereas the diffusional $^{22}\text{Na}^+$ uptake (0.5 min) and the equilibrium (120 min) uptake of $^{22}\text{Na}^+$ were not changed. The inhibition of Na^+ - H^+ exchange after ANF was likely due to alteration of the BBM antiporter itself, in that the H^+ conductance of BBMV was not increased. We conclude that synthetic ANF (a) decreases tubular Na^+ reabsorption linked to reabsorption of HCO_3^- in proximal tubules, and (b) inhibits proximal tubular reabsorption of P_i coupled to Na^+ reabsorption, independent of secretion and/or

action of parathyroid hormone or calcitonin. These ANF effects are associated with inhibition of Na^+ - P_i symport and of Na^+ - H^+ antiport in luminal BBMs. Our findings document that inhibition of Na^+ -coupled transport processes in proximal tubules is an integral part of the renal response to ANF.

Introduction

Recently, a family of polypeptides extracted from cardiac atrial tissue, designated as cardiopeptides or atrial natriuretic factor (ANF),¹ has been identified and subsequently chemically synthesized (1–4). The major known renal effects of these ANF peptides are natriuresis and changes in renal hemodynamics (2, 3). The natriuresis cannot be fully accounted for by changes in renal hemodynamics and/or physical factors governing in part tubular reabsorption of Na^+ (4), suggesting that ANF may inhibit directly the tubular transport of Na^+ . Unlike another recently described natriuretic factor (4), ANF has no effect on the activity of (Na^+ - K^+)-ATPase (4–6) or Na^+ transport in amphibian epithelia (5, 6).

Another partially related problem, is the question whether ANF inhibits tubular Na^+ reabsorption in the proximal or in the distal segments of the mammalian nephron. Although the bulk of the filtered Na^+ is reabsorbed in proximal tubules, a few initial studies seem to locate ANF inhibition of Na^+ reabsorption mainly in distal tubule segments (7, 8).

To address the question of whether or not ANF decreases tubular epithelial transport, namely in the proximal tubule, we studied the renal effects of infused synthetic rat ANF² peptide (9, 10).

The design of our study is based on the following premises. The filtered phosphate (P_i) is reabsorbed in proximal tubules via cotransport with Na^+ (11, 12), including secondary active Na^+ -dependent uptake of P_i across the luminal brush border membrane (BBM) (13, 14).

Na^+ is reabsorbed across the epithelium of the proximal tubule through several pathways (15). One of these important pathways for Na^+ reabsorption in the proximal tubule is a Na^+ cotransport with the bicarbonate anion (HCO_3^-) (15, 16). In this transport system, electroneutral Na^+ - H^+ antiport at the luminal BBM of proximal tubule (17) is the initial step in

1. *Abbreviations used in this paper:* ANF, atrial natriuretic factor; AO, acridine orange; BBM, brush border membrane; BBMV, brush border membrane vesicle; BP, blood pressure; FE, fractional excretion; GFR, glomerular filtration rate; P, plasma; PTH, parathyroid hormone; TPTX, thyroparathyroidectomy.

2. The atrial natriuretic factor used in this study is a synthetic polypeptide, which corresponds to the C-terminal polypeptide chain of 26 amino acid residues derived from the larger 73-amino acid precursor form of ANF (9), with sequence Arg-Arg-Ser-Ser-Cys-Phe-Gly-Gly-Arg-Ile-Asp-Arg-Ile-Gly-Ala-Gln-Ser-Gly-Leu-Gly-Cys-Asn-Ser-Phe-Arg-Tyr-COOH. A disulfide bridge links cysteines at positions 5 and 21 (9).

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Na⁺-linked reabsorption of HCO₃ and proximal tubular acidification (15, 16).

Therefore, if ANF inhibits Na⁺ reabsorption in proximal tubules, we can expect that Na⁺-P_i synport and Na⁺-H⁺ antiport across BBMs to be inhibited with a concomitant phosphaturia and bicarbonaturia.

With these considerations in mind, we studied the effects of the synthetic ANF (referred to as "ANF peptide" throughout the following text) on fractional excretion (FE) of Na⁺, P_i, HCO₃, and other electrolytes, as well as transport properties of the BBM vesicles (BBMVs) isolated from kidneys of the same animals.

Methods

Animals. Wistar rats of either sex weighing 200–225 g were fed standard rat diet containing 0.61% phosphorus (Ralston-Purina, Richmond, IN) ad libitum, with free access to water.

General experimental design. The studies were designed to minimize variability and to allow paired comparisons (18–20). To this end, animals that received infusion of ANF peptide and control animals that received vehicle were taken into the experiment on the same day, using the same set of solutions, biochemicals, and the same timing. Moreover, at the end of the clearance procedure, the BBMVs were prepared from kidneys of rats infused with ANF and from the controls at the same time, and transport measurements were also conducted simultaneously using the same radiochemicals, biochemicals, and standards. Therefore, glomerular filtration rate (GFR) and FE of electrolytes and subsequent transport studies on isolated BBMVs were determined simultaneously in two groups of rats.

Clearance experiments. On the day of the experiment the rats were anesthetized with Inactin (100 mg/kg) and prepared for clearance experiments. The animals were placed on a heated table, and body temperature was monitored with a rectal probe. A catheter was inserted into the carotid artery, and a basal blood sample was drawn. Acute thyroparathyroidectomy (TPTX) was performed by surgical removal of the glands and heat cautery. Adequacy of TPTX was confirmed in each group of rats by a fall in plasma calcium (ANF group: mean ± SEM, 2.50 ± 0.04 mM before and 2.35 ± 0.06 mM after TPTX, *n* = 7, *P* < 0.05; control group: 2.40 ± 0.05 mM before and 2.34 ± 0.04 mM after TPTX, *n* = 8, *P* < 0.05). A tracheostomy was performed, and the animals were allowed to breathe spontaneously. A catheter was inserted into a jugular vein for infusion of 2% inulin in normal saline at 3.5 ml/h. Urinary losses were replaced through a second jugular catheter. A bladder catheter was inserted. After TPTX a 2-h recovery period was allowed for attainment of a steady state. In the first group ("ANF group"), a control clearance was performed before intravenous infusion at 4.0 μg/kg per h of ANF (Merck, Sharp & Dohme, Rahway, NJ). After 20 min to obtain equilibrium of infused ANF, an experimental clearance was performed. In preliminary experiments, we found 4.0 μg/kg per h to be a maximally natriuretic nonhypotensive dose of ANF in the rat.

The second group shared the identical protocol except vehicle only—without ANF—was infused. Immediately after completion of the clearance studies, the kidneys were removed and decapsulated, and the cortex was rapidly dissected out and placed immediately in ice-cold solution (154 mM NaCl, 1 mM Tris-Hepes, pH 7.5) for preparation of a BBMv fraction (21).

BBMv transport studies. BBM fractions were prepared from homogenized renal cortical tissue by the slightly modified calcium precipitation procedure (2) described in full in our previous studies on rats (18–20, 22). The BBMVs were washed with 300 mM mannitol, 5 mM Tris-Hepes (pH 8.5), and were used immediately for solute uptake studies, for measurement of Na⁺-H⁺ exchange, or for testing of H⁺ efflux. All solutions used for the preparations of BBM fractions and for transport measurements were filtered through a 0.45-μm filter (Millipore Corp., Bedford, MA) on the day of use.

Uptake of [³²P]phosphate (18–20, 22) and L-[³H]proline (19, 22) by isolated BBMVs was measured by the Millipore filtration technique (18–20, 22). Briefly, BBMVs (~0.1 mg of protein) were suspended in a medium containing 100 mM mannitol, 100 mM NaCl, 5 mM Tris-Hepes (pH 8.5), and either 0.1 mM K₂H³²PO₄ or 0.025 mM L-[³H]proline (final concentrations). The uptake was measured at 20°C and was terminated at various times by addition of ice-cold 135 mM NaCl, 10 mM arsenate, 5 Tris-Hepes (pH 8.5). Uptake of ²²Na⁺ was determined in the absence of P_i or L-[³H]proline and by a similar procedure (20, 22) except that the incubation medium contained 100 mM mannitol, 100 mM NaCl, 5 mM Tris-Hepes (pH 8.5), and ²²NaCl (~4 × 10⁵ cpm/tube), and the uptake was terminated by the rapid addition of an ice-cold solution of 150 mM MgSO₄, buffered with 5 mM Tris-Hepes to pH 8.5, as in our previous study (22).

The rate of Na⁺-H⁺ exchange was determined by using a rapid filtration technique (23, 24), basically as described by Frierberg et al. (19), except that we employed BBMVs prepared by the calcium precipitation method, and that the pH of the intravesicular medium was pH 6.0 instead of 5.5. In the experiments when Na⁺-H⁺ exchange across the BBM was measured, BBMVs were prepared as in other solute uptake studies (18–20), but were preloaded (13) by resuspending and washing of the BBMv fraction in an acidic medium (23) that contains 150 mM KCl, 25 mM 2-(*N*-morpholino)-ethanesulfonic acid, 4.6 mM Tris, adjusted with KOH to pH 6.0. The rate of Na⁺-H⁺ exchange was measured by incubation of BBMVs preloaded with acidic intravesicular medium in a medium containing 144 mM KCl, 5 mM 2-(*N*-morpholino)-ethanesulfonic acid, 13 mM Tris, and 13 mM Hepes, adjusted with KOH to pH 7.5, and containing 1 mM ²²Na⁺Cl (0.1 μCi of ²²Na⁺ per tube) in final volume of 45 μl. The incubation was terminated by addition of 3 ml of ice-cold (0–2°C) "stop solution" containing 150 mM KCl, 16 mM Hepes, and 10 mM Tris, adjusted with KOH to pH 7.5 and rapid filtration (16), similarly as in other transport studies (12–15). Because the Na⁺-H⁺ antiport in the renal BBM is specifically blocked by amiloride (17), the ²²Na⁺ uptake at pH gradient condition (pH_i < pH_o) was always measured simultaneously with and without addition of 1 mM amiloride, the maximum inhibitory concentration of the compound (17). The amiloride-sensitive component of Na⁺-H⁺ exchange (~50%) was less than that reported in studies on rabbit (17, 25) and comparable to results of canine (26) BBMVs prepared by the Mg²⁺ precipitation procedure. This feature may be due to higher H⁺ conductance of rat BBMVs prepared by the Ca²⁺ precipitation method (26).

The rate of H⁺ efflux from BBMv was determined by using a method described by others (18, 20, 25, 26), which employs the quenching of acridine orange fluorescence as an indicator of the transmembrane proton gradient. Fluorescence was continuously monitored by using an AMINCO-Bowman spectrophotofluorometer (American Instrument Co., Silver Spring, MD) (excitation wavelength 493 nm, emission 530 nm) connected with a chart recorder. BBMVs were prepared at pH 6.0 as described above for Na⁺-H⁺ exchange measurements. The assay medium consisted of 6 μM acridine orange (AO) 150 mM KCl, 10 mM Tris-Hepes, pH 7.5. An aliquot (20 μl) of BBMVs (~150 μg of protein) equilibrated at pH 6.0 to the medium (final volume 2 ml) resulted in rapid quenching (about ~40%) of the AO fluorescence, which was followed by a slow increase in the fluorescence intensity as the imposed pH gradient dissipated. The addition of BBMVs equilibrated at pH 7.5 (H_i⁺ = H_o⁺) did not result in fluorescence quenching or in any time-dependent changes in fluorescence intensity. The rate of H⁺ efflux was expressed in relative units (18). The quenching of AO fluorescence after addition of BBMVs was taken as 100%. The rate of reappearance of the AO fluorescence was expressed as the relative (percent) rate of increase, for 15- and 30-s intervals (see Table VI).

In BBM transport studies (22–24, 27, 28), the final concentrations of radiolabeled solutes were 0.1 mM [³²P]_iphosphate, 0.025 mM L-[³H]proline, 0.05 mM [³H]glucose, and 100 mM ²²Na⁺Cl (21–23, 29). In most experiments that examined the ³²P_i uptake by BBMv, the uptake of L-[³H]proline was measured, for comparison, in the same aliquot of BBMv (22, 29).

The rates of uptake of $^{22}\text{Na}^+$ and other solutes, as well as the Na^+ - H^+ exchange are expressed in picomoles per milligram of BBMV protein per given time interval. In addition, the concentrative "uphill" Na^+ gradient-dependent uptake was also expressed in relative terms as a $\Delta\%$ ratio of solute uptake at the uphill phase to equilibrium phase (14, 21–23). In each BBM preparation, the uptake of solutes was measured in triplicate at each time period and the mean \pm SE was entered as $n = 1$. Uptake by BBMVs from each of the control and ANF groups was always compared on the same day to avoid interassay variations. Protein was determined by the method of Lowry et al. (30) after solubilization of the samples in 1% sodium dodecyl sulfate, as in our previous studies (21–23, 29).

Other analytic methods. Activities of BBM enzymes, i.e., alkaline phosphatase and leucine aminopeptidase, were assayed using methods previously employed and described in studies from this laboratory (28, 29). The specific activities of these enzymes were measured under conditions in which enzyme reaction rates were linear with respect to incubation time and protein concentrations. There was no difference in specific activity of alkaline phosphatase or leucine aminopeptidase between BBMs prepared from ANF-treated rats and from untreated controls.

Phosphate content of plasma (P_{P}) and urine was measured by the method of Chen et al. (31), inulin by the anthrone method (32), Na^+ and K^+ by flame photometry (23), and Ca and Mg by an atomic absorption spectrophotometer (Jarell-Ash Div., Fisher Scientific Co., Waltham, MA), as in previous studies (21, 22, 28, 29). Plasma bicarbonate was calculated from blood pH and PCO_2 measured on a blood gas analyzer (Instrumentation Laboratory Inc., Lexington, MA), and urine bicarbonate was calculated from urinary CO_2 content measured with a Natelson Microgasometer (Instrumentation Laboratory, Inc.), and urine pH (Instrumentation Laboratory, Inc.), as previously reported (33). Samples for urinary bicarbonate analysis were collected on mercury under oil. Plasma glucose was measured on a glucose oxidase analyzer (Beckman Instruments, Inc., Fullerton, CA).

Materials. Rats were supplied by Harlan Sprague Dawley, Inc. (Indianapolis, IN). $^{32}\text{P}_i$, L-[^3H]proline, and $^{22}\text{Na}^+$ were purchased from New England Nuclear (Boston, MA). Biochemicals, all of the highest purity grades available, were purchased from the Sigma Chemical Co. (St. Louis, MO) or other standard suppliers. Atrial natriuretic factor (ANF peptide) was a kind gift of Merck, Sharp & Dohme (Rahway, NJ). The ANF used was a polypeptide consisting of 26 amino acid residues² of the C-terminal of the precursor 73-amino acid form of ANF (9), synthesized by a combination of classical solution and solid-phase techniques as described by Seidah et al. (10). The peptide was stored in 50- μg aliquots in 0.05 ml of normal saline at -20°C .

Results

In the first series of experiments, we centered on the study of renal handling of P_i and BBM transport of P_i . Table I summarizes the clearance data on the effects of infusion of ANF, or vehicle in controls, on FE of electrolytes, GFR, blood pressure (BP), and plasma parameters in TPTX rats from which subsequently BBMVs were isolated for BBM transport studies of P_i in vitro. Infusion of ANF was associated, on the one hand, with significant increases in FE_{P_i} , FE_{Na} , and GFR both when compared with the control period (first period) in the same animal, and also by paired analysis, i.e., compared with the matched control rats studied simultaneously. On the other hand, FE_{Ca} , FE_{Mg} , FE_{K} , or BP and P_{P} did not change when ANF was infused (Table I). Plasma glucose did not change with ANF infusion (137 ± 5 ng/dl, mean \pm SEM in the first control period and 128 ± 8 ng/dl, $n = 7$, during ANF infusion, second period). In the control rats infused with vehicle only, there were no changes in GFR or FE of any of the electrolytes, BP, or P_{P} (Table I).

Table II summarizes the Na^+ gradient-dependent uptake

Table I. Effects of Infusion of ANF on Excretion of Electrolytes and Other Parameters in Clearance Studies in TPTX Rats Subsequently Used for Measurement of Transport of Phosphate, L-Proline, and Na^+ by BBMVs

	First period (vehicle)	Second period ^{II}
ANF-infused group ($n = 7$)*		
FE_{P_i} (%)	3.51 ± 1.49	$9.64\pm 1.97\ddagger$
P_{P} (mM)	2.46 ± 0.19	2.13 ± 0.19
FE_{Na} (%)	1.30 ± 0.26	$4.65\pm 0.58\ddagger$
FE_{Ca} (%)	3.20 ± 0.7	3.70 ± 0.5
FE_{K} (%)	37.80 ± 8.6	36.20 ± 5.5
FE_{Mg} (%)	16.30 ± 4.2	20.30 ± 5.4
GFR (ml/min)	2.17 ± 0.31	$3.63\pm 0.39\ddagger$
BP (mmHg)	131.00 ± 5.0	131.00 ± 5
Control vehicle-infused group ($n = 7$)*		
FE_{P_i} (%)	1.61 ± 1.09	1.93 ± 1.02
P_{P} (mM)	2.21 ± 0.15	2.44 ± 0.18
FE_{Na} (%)	1.01 ± 0.29	2.27 ± 1.58
FE_{Ca} (%)	2.90 ± 0.6	4.10 ± 1.3
FE_{K} (%)	32.20 ± 9.2	26.20 ± 3.5
FE_{Mg} (%)	17.50 ± 4.6	24.10 ± 5.9
GFR (ml/min)	3.56 ± 0.48	2.63 ± 0.58
BP (mmHg)	130.00 ± 3.0	128.00 ± 3.0

* n denotes number of animals; ANF-infused animals were paired with controls.

\ddagger $P < 0.05$ compared with control by paired t test (\ddagger) or ANF with control groups by unpaired t test (\S).

^{II} ANF-infused group infused with ANF in the second period; control group infused with vehicle in the second period.

of [^{32}P]phosphate and L-[^3H]proline by BBMVs from ANF-infused rats and vehicle-infused control rats. Infusion of ANF decreased the concentrative Na^+ gradient-dependent BBMVs uptake of $^{32}\text{P}_i$ both in absolute terms, i.e., net rate (picomoles per milligram of protein) and relative terms ($\Delta\%$ overshoot) both at 15 and 30 s. In contrast, the $^{32}\text{P}_i$ uptake at the equilibrium phase (120 min) was not different between BBMVs from controls and ANF-infused rats. In aliquots of the same BBMVs preparations, the concentrative Na^+ gradient-dependent uptake of L-[^3H]proline was not different when BBMVs from ANF-infused rats were compared with controls (Table II, Fig. 1), and also, the uptake at 120 min was identical (Table II).

Likewise, the Na^+ gradient-dependent uptake of D-[^3H]glucose (expressed in picomoles per milligram of protein per 15 s; mean \pm SEM) in BBMVs from ANF-treated rats (588 ± 64 ; $n = 4$) was not significantly different from the D-[^3H]glucose uptake in BBMVs from control rats (659 ± 35 ; $n = 4$).

The diffusional uptake of $^{22}\text{Na}^+$ ($\text{Na}_0^+ > \text{Na}_i^+$; $\text{pH}_0 = \text{pH}_i$), measured in the absence of a proton gradient (intravesicular pH equal to extravesicular pH), was not different between BBMVs from control and from ANF-infused animals, either at 30 s or after equilibrium at 100 min (Table III). The decrease in Na^+ gradient-dependent uptake of $^{32}\text{P}_i$ at 30 s by BBMVs was significantly correlated with the increase in FE_{P_i} determined in the same animal ($r = 0.99$; $P < 0.001$; t test, $n = 7$), but not with FE_{K} , FE_{Ca} , or FE_{Mg} . In contrast, the FE_{P_i}

Table II. Effects of ANF Infusion on BBMV Transport of $^{32}\text{P}_i$ and L-[^3H]Proline (See Table I)

Time	Controls (n = 7)	ANF-treated (n = 7)
	pmol/mg protein	pmol/mg protein
[$^{32}\text{P}_i$]Phosphate uptake		
15 s	1,862±69*	1,378±35‡
30 s	2,286±152	1,755±143‡
120 min	803±56	752±66
% $\Delta_{15\text{s}}$ §	150±22	102±21‡
% $\Delta_{30\text{s}}$ §	194±29	141±22‡
L-[^3H]Proline uptake		
15 s	286±33*	280±29
120 min	41±4	38±4
% $\Delta_{15\text{s}}$ §	607±103	644±86

* Mean±SEM of seven experiments; controls and ANF-treated were paired.

‡ Significantly different from corresponding control values ($P < 0.05$) by paired (*) or group (§) t test.

§ Ratio of initial uptake ("overshoot") expressed relative to equilibrium point (120 min).

did not correlate with the rate of $^{22}\text{Na}^+$ uptake by BBMVs at 30 s, or with uptake of L-[^3H]proline.

The effect of direct addition of ANF in vitro on P_i transport was measured in BBMVs prepared from nontreated rats. BBMVs were preincubated with ANF (4 ng/ml in a Na^+ -free buffer) in vitro for 30 s at 20°C and then the Na gradient-

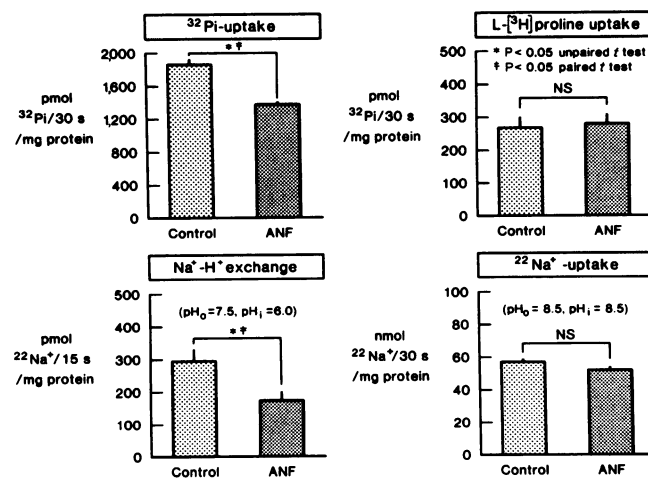


Figure 1. Transport properties of renal cortical BBMVs from rats infused with vehicle (control) and rats infused with ANF peptide (ANF). The Na^+ -dependent uptake of P_i and L-[^3H]proline, as well as the uptake of $^{22}\text{Na}^+$ and amiloride-sensitive Na^+-H^+ exchange were measured in the initial (15 s, 30 s) time periods. Each bar denotes mean±SE of paired experiments ($^{32}\text{P}_i$ and L-[^3H]proline, $n = 7$; $^{22}\text{Na}^+$ uptake, $n = 6$; amiloride-sensitive Na^+-H^+ exchange, $n = 4$). Asterisk (*) denotes statistically significant difference from controls based on group comparison ($P < 0.05$, unpaired t test). Double dagger (§) denotes statistically significant difference from controls based on paired comparison (paired t test, $P < 0.05$).

Table III. Uptake of $^{22}\text{Na}^+$ in the Absence of pH Gradient ($\text{pH}_0 = \text{pH}_i$; $\text{Na}_0^+ = 100 \text{ mM}$; $\text{Na}_i^+ = 0$) by BBMV from Control and from ANF-treated Rats

Animals	$^{22}\text{Na}^+$ uptake	
	30 s	120 min
	nmol/mg protein	nmol/mg protein
Controls	57±2*	185±11
ANF-treated	52±2	166±10

* All values are mean±SEM of six experiments. Uptake of $^{22}\text{Na}^+$ in BBMVs from ANF-infused rats was not significantly different from controls (t test).

dependent uptake of $^{32}\text{P}_i$ was measured after 30 s, as described in Methods. Under these experimental conditions, the rate of P_i uptake was not changed by the presence of ANF (data not shown).

In the next series of experiments, we studied the effect of ANF on HCO_3^- excretion and Na^+-H^+ antiport in BBMVs. Table IV summarizes the clearance data and Table V, the analysis of the rate of Na^+-H^+ exchange across BBMVs

Table IV. Effects of Infusion of ANF on Excretion of Electrolytes and Other Parameters in TPTX Rats Subsequently Used for Measurements of Na^+-H^+ Exchange in BBMV Studies (See Table V)

	First period (vehicle)	Second period ^{II}
ANF-infused group (n = 8)*		
$\text{FE}_{\text{HCO}_3^-}$ (%)	5.87±2.14	9.99±1.99‡§
FE_{Na^+} (%)	1.06±0.26	3.73±0.72‡§
FE_{P_i} (%)	0.84±0.41	4.99±1.97‡§
$\text{FE}_{\text{Ca}^{2+}}$ (%)	2.00±0.6	2.80±0.6
FE_{K^+} (%)	26.80±2.9	30.00±5.0
GFR (ml/min)	2.92±0.27	4.41±0.53‡§
BP (mmHg)	129.00±5.0	130.00±5.0
P_{PO_4} (mM)	2.66±0.07	2.97±0.20
$\text{P}_{\text{HCO}_3^-}$ (mM)	27.00±1.00	27.00±1.00
Control vehicle-infused group (n = 8)*		
$\text{FE}_{\text{HCO}_3^-}$ (%)	5.62±1.38	6.09±1.60
FE_{Na^+} (%)	1.17±0.24	0.92±0.26
FE_{P_i} (%)	0.69±0.25	0.90±0.38
FE_{K^+} (%)	28.30±6.2	23.40±5.4
GFR (ml/min)	2.88±0.38	2.72±0.47
BP (mmHg)	130.00±4.0	129.00±5.0
P_{PO_4} (mM)	3.04±0.17	2.97±0.15
$\text{P}_{\text{HCO}_3^-}$ (mM)	27.00±1.00	27.00±1.00

* n denotes number of animals; controls and ANF-infused animals were paired.

‡§ $P < 0.05$ compared with control by paired t test (§) or ANF with control groups by unpaired t test (§).

^{II} ANF-infused group infused with ANF in the second period; control group infused with vehicle in the second period.

Table V. Effect of ANF Infusion on the Rate of $\text{Na}^+ - \text{H}^+$ Exchange across BBMVs in the Presence of Proton Gradient [$\text{H}_i^+ > \text{H}_o^+$]

	$^{22}\text{Na}^+$ uptake	
	15 s	120 min
Controls		
– Amiloride	539±33*	1,275±66
+ Amiloride	244±11	1,430±81
Amiloride-sensitive	295±36	—
ANF-treated		
– Amiloride	448±26‡§	1,222±63
+ Amiloride	275±5‡§	1,346±98
Amiloride-sensitive	173±28‡§	—

$^{22}\text{Na}^+$ uptake was measured with or without added amiloride. The “amiloride-sensitive” uptake was determined as a differential of uptake in the presence and in the absence of 1 mM amiloride, measured in the same BBMV preparation.

* All values are mean±SEM of four experiments (pooled tissue from eight rats).

‡§ Significantly different from corresponding control values ($P < 0.05$ or higher degree of significance; paired t test(‡), group t test(§).

prepared from the same animals. As in the first series of experiments (Table I), the infusion of ANF was associated with significant increases in FE_{Na} , FE_{P_i} , and GFR. The FE_{HCO_3} was markedly increased after ANF infusion (second period) both when compared with the first control period in the same animals, or when compared with the control group receiving vehicle only in the second period (Table IV). Again, as in previous experiments, the FE of Ca^{2+} and K^+ , BP , and P_i and P_{HCO_3} did not change when ANF was infused (Table IV). Likewise, the pH and PCO_2 did not change in response to ANF (data not shown). In the control group (Table IV), there was no change in GFR, FE of any electrolytes, BP, or P_i .

Table V summarizes the effects of ANF infusion on the rate of $\text{Na}^+ - \text{H}^+$ exchange across BBMVs in the presence of a proton gradient. The total $^{22}\text{Na}^+$ uptake rate (15 s) in the presence of H^+ gradient ($\text{H}_i^+ > \text{H}_o^+$) was significantly lower (–17%) in BBMVs for ANF-infused rats. The difference between BBMVs from control and ANF-infused rats was even more pronounced (–41%) in the amiloride-sensitive component of $\text{Na}^+ - \text{H}^+$ exchange across BBM (Table V). After equilibration (120 min) the $^{22}\text{Na}^+$ uptake did not differ between control and ANF-infused rats (Table V).

The rate of H^+ efflux from BBMVs [$\text{H}_i^+ > \text{H}_o^+$] in the absence of Na^+ was determined using the quenching and subsequent reappearance of AO fluorescence as an indicator of the transmembrane proton-gradient dissipation, is shown in Table VI. The rates of relative increase in the AO fluorescence intensity at 15- and 30-s intervals measured under similar conditions for control and ANF-treated vesicles were not different. These results indicate that the H^+ conductance of the BBM was not increased, and hence, cannot account for decreased $\text{Na}^+ - \text{H}^+$ exchange rate in BBMVs from ANF-treated rats.

To explore the relationship between changes in excretions of Na^+ and other electrolytes elicited by ANF, we examined

Table VI. Effect of ANF Administration In Vivo on the Rate of Dissipation of Proton Gradient [$\text{H}_i^+ > \text{H}_o^+$] across BBMVs

Experiment	Condition	+Δ% Fluorescence	
		15 s	30 s
1	Controls	17.3±0.5*	30.0±1.1
	ANF-treated	17.4±0.4	31.2±1.1
2	Controls	20.0±1.1	34.6±1.1
	ANF-treated	20.2±0.5	36.4±1.2

The proton efflux was determined as a rate of return of AO fluorescence in relative (%) units. For further details, see Methods.

* Each number denotes mean±SEM of five measurements.

correlations between changes (Δ) in FE_{Na} and FE of other simultaneously measured ions. Table VII summarizes the correlation of change in FE of Na^+ ($\Delta\text{FE}_{\text{Na}}$) with the change in FE of phosphate ($\Delta\text{FE}_{\text{P}_i}$), bicarbonate ($\Delta\text{FE}_{\text{HCO}_3}$), potassium ($\Delta\text{FE}_{\text{K}}$), magnesium ($\Delta\text{FE}_{\text{Mg}}$), and calcium ($\Delta\text{FE}_{\text{Ca}}$). Results show that $\Delta\text{FE}_{\text{Na}}$ correlated significantly with $\Delta\text{FE}_{\text{P}_i}$ and $\Delta\text{FE}_{\text{HCO}_3}$, but not with $\Delta\text{FE}_{\text{K}}$, $\Delta\text{FE}_{\text{Mg}}$, or $\Delta\text{FE}_{\text{Ca}}$.

Discussion

As expected from previous reports (1, 4), infusion of ANF peptide significantly increased the FE_{Na} in TPTX rats. Even more importantly, ANF infusion also significantly decreased reabsorption of P_i and HCO_3 anions, the bulk of which are reabsorbed from glomerular filtrate in proximal tubules. As in other studies (1–5), infusion of ANF increases GFR, and hence, filtered loads of solutes, including Na^+ , HCO_3 , and P_i . When electrolyte excretions are expressed as FE relative to filtered amount, it should be realized that net tubular reabsorption of solutes is increased. However, in response to ANF, the amount of reabsorbed Na^+ , HCO_3 , and P_i is less than proportional to filtered loads due to increased GFR and such phenomenon indicates that tubular reabsorptive transport is inhibited by ANF. The decreased reabsorptions of P_i and HCO_3 were specific because the FE of Ca^{2+} , K^+ , and Mg^{2+} were not affected. In that our experiments were conducted in TPTX rats, the effects of ANF peptide cannot be explained by an effect on the release or potentiation of action of endogenous

Table VII. Correlation between Changes in Fractional Excretion of Sodium ($\Delta\text{FE}_{\text{Na}}$) and Changes in Fractional Excretions (ΔFE) of Other Measured Electrolytes

Change	r^*	n^\ddagger	P value§
$\Delta\text{FE}_{\text{P}_i}$	0.90	7	<0.01
$\Delta\text{FE}_{\text{HCO}_3}$	0.89	7	<0.01
$\Delta\text{FE}_{\text{K}}$	0.71	7	NS
$\Delta\text{FE}_{\text{Ca}}$	0.62	7	NS
$\Delta\text{FE}_{\text{Mg}}$	0.60	7	NS

* Correlation coefficient.

‡ Number of pairs.

§ For significance of r , t test.

parathyroid hormone (PTH) or calcitonin, polypeptide hormones that are well known to increase renal excretion of P_i and HCO_3^- (34), or by removal of the antiphosphaturic effects of thyroid hormones (22).

ANF-elicited changes in the FE_{Na} showed a strong positive correlation with changes in FE_{P_i} and $FE_{HCO_3^-}$, whereas no significant correlation was noted with FE_{Ca} , FE_{Mg} , or FE_K (Table VII). These correlative relationships further suggest that ANF peptide inhibited reabsorption of Na^+ , which is linked to proximal tubular reabsorption of P_i and HCO_3^- . This suggestion is strongly supported by findings of changed transport properties of BBMVs isolated from the same rats. Of particular interest is a finding that Na^+ gradient-dependent uptake of P_i was markedly inhibited in response to infusion of ANF peptide (Table II, Fig. 1). A lack of any effect of the infusion of ANF peptide on the (30 s) diffusional $^{22}Na^+$ uptake by BBMVs (Table III), or on other transport systems energized by ($Na_0^+ > Na_i^+$) gradient, i.e., the uptake of L-[3H]proline and D-[3H]glucose indicate that the inhibition elicited by infusion of ANF peptide is relatively specific for the BBM transport system for P_i (Table III, Fig. 1). Therefore, the decrease in the Na^+ -dependent P_i uptake is not due to increased diffusional permeability of BBM for Na^+ , hence the faster dissipation of ($Na_0^+ > Na_i^+$) gradient in response to the infused ANF peptide (35). Furthermore, the observations that BBMVs uptake of L-[3H]proline and of $^{22}Na^+$ after complete equilibration (120 min) (Tables II and III) is not different after the infusion of ANF peptide indicate that intravesicular BBMVs volume is not altered (14, 21). Therefore, the observed decrease in Na^+ gradient-dependent BBMVs uptake of P_i provides a strong argument for the notion that proximal tubular P_i reabsorption was reduced in response to ANF. As in the case of other hormones that modulate BBM transport of P_i in vivo but not in vitro, e.g., PTH (35a), glucocorticoids (28), or T_4 (22) direct interaction of ANF with the BBM in vitro had no effect. This indicates that ANF requires other cellular components and/or intact cells for expression of its effect on the BBM transport system.

The Na^+ - H^+ antiport at the BBM, a major component of HCO_3^- -linked proximal tubular Na^+ reabsorption (11, 15–17, 36), was markedly blunted by ANF-peptide infusion (Table V). This decreased capacity of the Na^+ - H^+ exchange mechanism in the BBM may be the basis of diminished Na^+ -linked HCO_3^- reabsorption in proximal tubules, which in turn increases $FE_{HCO_3^-}$ (Table IV) (15, 16, 36, 37). A finding that ANF infusion does not lead to change in H^+ conductance of the BBM, and hence, in the rate of dissipation of [$H_i^+ > H_0^+$] gradient, suggests that alteration of Na^+/H^+ antiporter within the BBM is a basis of lower rate of Na^+ luminal entry in exchange to H^+ (25, 26). Besides luminal entry across BBM, another major component operative in transepithelial Na^+ reabsorption and in Na^+ -linked transport systems in proximal tubules is the activity of (Na^+ - K^+)-ATPase at basolateral membranes, an active Na^+ pump that continuously maintains low intracellular Na^+ and lumen-to-cell Na^+ gradient (11–15). Studies reported by other investigators show that ANF does not influence (Na^+ - K^+)-ATPase activity (4–6).

Taken together, the observations that the rate of Na^+ - H^+ antiport and Na^+ - P_i synport of BBM was diminished (Fig. 1) indicate that ANF-peptide inhibited at least in part the Na^+ reabsorption and reabsorption of solutes cotransported with Na^+ in proximal tubules (19, 20, 38–40). It remains to be further explored whether or not the changes in BBM transport

consequent to ANF infusion are the only and/or the primary basis of altered urinary electrolyte excretion.

Major qualifications and potential limitations in the interpretation of the present observations should be briefly mentioned. First, the ANF peptide was infused in vivo, and it is not possible at the present time to conclude whether the changes in renal function (ΔFE) and the BBM effects were due to the direct action of this peptide on the proximal tubular cells or whether ANF acts through some indirect mechanism. In that ANF does not inhibit transport by direct addition in vitro, it is unlikely to interact with BBM transporters directly and requires the whole cell for its action. Conceivably, the ANF peptide can also be biotransformed in vivo to the substance, e.g., the proteolytic peptide fragment, that may be the compound directly interacting with proximal tubules. Also, ANF administration may affect release of endogenous factors that could influence the renal response, such as aldosterone (41). PTH, calcitonin, and glucagon can be excluded based on the experimental design—TPTX rats and findings of the lack of changes in the glucose level. Moreover, the possibility that ANF would act on tubular receptors for PTH or calcitonin or glucagon is unlikely, because the ANF-peptide does not share an homologous amino acid sequence with any other known biologically active peptide (42). The time course and reversibility of the infused ANF peptide on BBM transport systems remain to be explored. Second, the substantial increases in GFR (Tables I and IV) may contribute to the increases in FE of electrolytes. It is highly unlikely that the entire effect is due to hemodynamic changes, as indicated by recent study on anesthetized dogs (43).

It is a most plausible explanation that the diminished luminal entry of P_i with Na^+ and in exchanges with H^+ account for the lower rate of proximal tubule reabsorption of these solutes.

In summary, the present observations indicate that ANF peptide (and possibly other derivatives of this peptide hormone family) cause phosphaturia and bicarbonaturia, and inhibit the Na^+ -linked transport process in the proximal segments of the tubules. Therefore, the effect of ANF on proximal tubular transport of solutes constitutes another component of renal actions (1, 4) of this intriguing, newly found regulatory substance.

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